

STIC-Biotech/Ch mLib

85933

From: Schultz, James  
Sent: Tuesday, February 04, 2003 10:18 AM  
To: STIC-Biotech/ChemLib  
Subject: Sequence search 09/960,143

12E18

Hello,  
Could you please run a length limited nucleotide sequence search on SEQ ID NO:3 (1639 nt long) in the above entitled application, where the maximum size of the returned hits are no longer than 50 nucleotides?  
Thanks,  
Doug Schultz

J. Douglas Schultz, Ph.D.  
AU 1635 (Biotechnology)  
Patent Examiner  
United States Patent and Trademark Office  
(703) 308-9355  
(703) 746-3973 (fax)  
Office: CM1 12E18  
Mail: CM1 11E12

CARE

Point of Contact:  
Thomas G. Larson, Ph.D.  
703-308-7309  
CM1, Rm. 6 B 01

Searcher: Larson  
Phone: \_\_\_\_\_  
Location: \_\_\_\_\_  
Date Picked Up: 2/4  
Date Completed: 2/13  
Searcher Prep/Review: 10  
Clerical: \_\_\_\_\_  
Online time: 5

TYPE OF SEARCH:  
NA Sequences: 1  
AA Sequences: \_\_\_\_\_  
Structures: \_\_\_\_\_  
Bibliographic: \_\_\_\_\_  
Litigation: \_\_\_\_\_  
Full text: \_\_\_\_\_  
Patent Family: \_\_\_\_\_  
Other: \_\_\_\_\_

VENDOR/COST (where applic.)  
STN: \_\_\_\_\_  
DIALOG: \_\_\_\_\_  
Questel/Orbit: \_\_\_\_\_  
DRLink: \_\_\_\_\_  
Lexis/Nexis: \_\_\_\_\_  
Sequence Sys.: ABSS03  
WWW/Internet: \_\_\_\_\_  
Other (specify): \_\_\_\_\_

**THIS PAGE BLANK (USPTO)**

Result No.	Score	Query %		Length	DB	ID	Description
		Match					
1	37.4	2.3	47	1	US-08-330-163-42	Sequence 42, Appl	
2	37.4	2.3	47	1	US-08-482-111-42	Sequence 42, Appl	
C 3	37	2.3	37	1	US-08-330-163-40	Sequence 40, Appl	
C 4	37	2.3	37	1	US-08-482-111-40	Sequence 40, Appl	
C 5	33	2.0	33	2	US-08-410-654B-42	Sequence 42, Appl	
C 6	33	2.0	33	2	US-08-474-851-42	Sequence 42, Appl	
C 7	33	2.0	33	2	US-08-481-560-42	Sequence 42, Appl	
C 8	31	1.9	31	5	PCT-US96-08142-9	Sequence 9, Appl	
C 9	30	1.8	30	4	US-09-368-160B-6	Sequence 6, Appl	
C 10	30	1.8	30	4	US-09-456-399-6	Sequence 6, Appl	
C 11	28	1.7	28	2	US-08-859-998-479	Sequence 479, Appl	
C 12	28	1.7	28	2	US-08-859-998-480	Sequence 480, Appl	
C 13	28	1.7	28	4	US-09-235-928-479	Sequence 479, Appl	
C 14	28	1.7	28	4	US-09-235-928-480	Sequence 480, Appl	
C 15	25	1.5	25	1	US-08-327-494A-8	Sequence 8, Appl	
C 16	25	1.5	25	5	PCT-US95-13659-8	Sequence 8, Appl	
C 17	25	1.5	25	5	PCT-US96-08142-8	Sequence 8, Appl	
C 18	24	1.5	24	4	US-09-710-200-72	Sequence 72, Appl	
C 19	24	1.5	24	5	PCT-US96-08142-7	Sequence 72, Appl	
C 20	23.4	1.4	47	4	US-09-641-638-1167	Sequence 1167, Appl	
C 21	23	1.4	23	5	US-08-327-494A-7	Sequence 7, Appl	
C 22	23	1.4	23	5	PCT-US95-13659-7	Sequence 7, Appl	
C 23	23	1.4	40	1	US-08-482-111-53	Sequence 53, Appl	
C 24	22	1.3	22	4	US-09-046-894-20	Sequence 20, Appl	
C 25	22	1.3	48	1	US-08-317-102-1	Sequence 1, Appl	
C 26	21.8	1.3	47	4	US-09-338-907-250	Sequence 250, Appl	
C 27	21.8	1.3	47	4	US-09-338-907-327	Sequence 327, Appl	

RESULT 3  
 US-08-330-163-40/c  
 Sequence to Application US/08330163  
 Patent No. 565672A  
 GENERAL INFORMATION:  
 APPLICANT: Daly, Thomas J.  
 APPLICANT: Larosa, Gregory J.  
 TITLE OF INVENTION: Chemokine-Like Proteins and Methods of  
 TITLE OF INVENTION: Use  
 NUMBER OF SEQUENCES: 46  
 CORRESPONDENCE ADDRESS:  
 ADDRESSEE: Fish & Richardson  
 STREET: 225 Franklin Street  
 CITY: Boston  
 STATE: MA  
 COUNTRY: U.S.A.  
 ZIP: 02110-2804  
 COMPUTER READABLE FORM:  
 MEDIUM TYPE: Floppy disk  
 COMPUTER: IBM PC compatible  
 OPERATING SYSTEM: PC-DOS/MS-DOS

Query Match 2.38; Score 37; DB 1; Length 37;

Best Local Similarity 100.0%; Pred. No. 18;  
Matches 37; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 351 GGTGTGGAGAGCTTTTGAAGAGGGCTGAGATTCA 387  
Db 37 GGTGTGGAGAGCTTTTGAAGAGGGCTGAGATTCA 1

RESULT 5  
US-08-410-654B-42/c  
Sequence 42, Application US/08410654B  
Patent No. 5833976  
GENERAL INFORMATION:  
APPLICANT: Rene de Waal Malefyt  
APPLICANT: Di-Hwei Hsu  
APPLICANT: Anne O'Garra  
APPLICANT: Hergen Spits  
TITLE OF INVENTION: Use of Interleukin-10 to Treat  
TITLE OF INVENTION: Septic Shock  
NUMBER OF SEQUENCES: 61  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Schering-Plough Corporation  
STREET: 2000 Galloping Hill Road  
CITY: Kenilworth  
STATE: New Jersey  
COUNTRY: USA  
ZIP: 07033

COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: Macintosh  
OPERATING SYSTEM: 7.5.3  
SOFTWARE: Microsoft Word 5.1a  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/410,654B  
FILING DATE: 24-MAR-1995  
CLASSIFICATION: 424  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/229,854  
FILING DATE: 19-APR-1994  
APPLICATION NUMBER: US 07/926,853  
FILING DATE: 06-AUG-1992  
APPLICATION NUMBER: US 07/742,129  
FILING DATE: 06-AUG-1991  
ATTORNEY/AGENT INFORMATION:  
NAME: Foulke, Cynthia L.  
REGISTRATION NUMBER: 32,364  
REFERENCE/DOCKET NUMBER: DX0221KQ1  
TELEPHONE: 908-298-2987  
TELEFAX: 908-298-5388  
INFORMATION FOR SEQ ID NO: 42:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 33 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: double  
TOPOLOGY: linear  
MOLECULE TYPE: DNA (oligonucleotide)  
US-08-410-654B-42

Query Match 2.0%; Score 33; DB 2; Length 33;  
Best Local Similarity 100.0%; Pred. No. 1.1e+02;  
Matches 33; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 358 GAGAAGTTTTTGAAGAGGGCTGAGAATTCATAA 390  
Db 33 GAGAAGTTTTTGAAGAGGGCTGAGAATTCATAA 1

RESULT 5  
US-08-410-851-42/c  
Sequence 42, Application US/08474851  
Patent No. 5837232  
GENERAL INFORMATION:  
APPLICANT: Rene de Waal Malefyt  
APPLICANT: Di-Hwei Hsu  
APPLICANT: Anne O'Garra  
APPLICANT: Hergen Spits  
TITLE OF INVENTION: Use of Interleukin-10 to Modulate  
TITLE OF INVENTION: Inflammation or T-Cell Mediated  
NUMBER OF SEQUENCES: 61  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Schering-Plough Corporation  
STREET: 2000 Galloping Hill Road

APPLICANT: Rene de Waal Malefyt  
APPLICANT: Di-Hwei Hsu  
APPLICANT: Anne O'Garra  
APPLICANT: Hergen Spits  
TITLE OF INVENTION: Use of An Interleukin-10 Antagonist to Treat  
TITLE OF INVENTION: A B Cell Mediated Autoimmune Disorder  
NUMBER OF SEQUENCES: 61  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Schering-Plough Corporation  
STREET: 2000 Galloping Hill Road  
CITY: Kenilworth  
STATE: New Jersey  
COUNTRY: USA  
ZIP: 07033

COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: Macintosh  
OPERATING SYSTEM: 7.5.3  
SOFTWARE: Microsoft Word 6.0  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/474,851  
FILING DATE: 07-JUN-1995  
CLASSIFICATION: 424  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: 08/410,654  
FILING DATE: 24-MAR-1995  
APPLICATION NUMBER: US 08/229,854  
FILING DATE: 19-APR-1994  
APPLICATION NUMBER: US 07/926,853  
FILING DATE: 06-AUG-1992  
APPLICATION NUMBER: US 07/742,129  
FILING DATE: 06-AUG-1991  
ATTORNEY/AGENT INFORMATION:  
NAME: Foulke, Cynthia L.  
REGISTRATION NUMBER: 32,364  
REFERENCE/DOCKET NUMBER: DX0221KQ1GD  
TELEPHONE: 908-298-2987  
TELEFAX: 908-298-5388  
INFORMATION FOR SEQ ID NO: 42:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 33 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: double  
TOPOLOGY: linear  
MOLECULE TYPE: DNA (oligonucleotide)  
US-08-474-851-42

Query Match 2.0%; Score 33; DB 2; Length 33;  
Best Local Similarity 100.0%; Pred. No. 1.1e+02;  
Matches 33; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 358 GAGAAGTTTTTGAAGAGGGCTGAGAATTCATAA 390  
Db 33 GAGAAGTTTTTGAAGAGGGCTGAGAATTCATAA 1

RESULT 7  
US-08-481-560-42/c  
Sequence 42, Application US/08481560  
Patent No. 5837293  
GENERAL INFORMATION:  
APPLICANT: Rene de Waal Malefyt  
APPLICANT: Di-Hwei Hsu  
APPLICANT: Anne O'Garra  
APPLICANT: Hergen Spits  
TITLE OF INVENTION: Use of Interleukin-10 to Modulate  
TITLE OF INVENTION: Inflammation or T-Cell Mediated  
NUMBER OF SEQUENCES: 61  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Schering-Plough Corporation  
STREET: 2000 Galloping Hill Road

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; CITY: Kenilworth
; STATE: New Jersey
; COUNTRY: USA
; ZIP: 07033
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: Macintosh
; OPERATING SYSTEM: 7.5.3
; SOFTWARE: Microsoft Word 6.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/481.560
; FILING DATE: 07-JUN-1995
; CLASSIFICATION: 424
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/410,654
; FILING DATE: 24-MAR-1995
; APPLICATION NUMBER: US 08/229,854
; FILING DATE: 19-APR-1994
; APPLICATION NUMBER: US 07/926,853
; FILING DATE: 06-AUG-1992
; APPLICATION NUMBER: US 07/742,129
; FILING DATE: 06-AUG-1991
; ATTORNEY/AGENT INFORMATION:
; NAME: Foulke, Cynthia L.
; REGISTRATION NUMBER: 32,364
; REFERENCE/DOCKET NUMBER: DX0221KQIGC
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 908-298-2987
; TELEFAX: 908-298-5388
; INFORMATION FOR SEQ ID NO: 42:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 33 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (oligonucleotide)
; US-08-481-560-42

Query Match 2.0%; Score 33; DB 2; Length 33;
Best Local Similarity 100.0%; Pred. No. 1.1e+02; Indels 0; Gaps 0;
Matches 33; Conservative 0; Mismatches 0;

QY 358 GAGAAATTTTGAAGAGGGCTGAGAATTCATAA 390
Db 33 GAGAAATTTTGAAGAGGGCTGAGAATTCATAA 1

RESULT 8
PCT-US96-08142-9/c
; Sequence 9, Application PC/TUS9608142
; GENERAL INFORMATION:
; APPLICANT: Constance Emmett, Kimberly A. Foster
; TITLE OF INVENTION: Universal Chemistry Enzyme-Linked
; TITLE OF INVENTION: Immunosorbent Assay for Detection of
; TITLE OF INVENTION: mRNA Expression
; NUMBER OF SEQUENCES: 9
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Jane Massey Licata, Esq.
; STREET: 210 Lake Drive East, Suite 201
; CITY: Cherry Hill
; STATE: NJ
; COUNTRY: USA
; ZIP: 08002
; COMPUTER READABLE FORM:
; MEDIUM TYPE: DISKETTE, 3.5 INCH, 1.44 Mb STORAGE
; COMPUTER: IBM 486
; OPERATING SYSTEM: WINDOWS FOR WORKGROUPS
; SOFTWARE: WORDPERFECT 5.1
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: PCT/US96/08142
; FILING DATE: Herewith
; CLASSIFICATION:
; PRIOR APPLICATION DATA:

; CITY: Kenilworth
; STATE: New Jersey
; COUNTRY: USA
; ZIP: 07033
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: Macintosh
; OPERATING SYSTEM: 7.5.3
; SOFTWARE: Microsoft Word 6.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/481.560
; FILING DATE: 07-JUN-1995
; CLASSIFICATION: 424
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/410,654
; FILING DATE: 24-MAR-1995
; APPLICATION NUMBER: US 08/229,854
; FILING DATE: 19-APR-1994
; APPLICATION NUMBER: US 07/926,853
; FILING DATE: 06-AUG-1992
; APPLICATION NUMBER: US 07/742,129
; FILING DATE: 06-AUG-1991
; ATTORNEY/AGENT INFORMATION:
; NAME: Foulke, Cynthia L.
; REGISTRATION NUMBER: 32,364
; REFERENCE/DOCKET NUMBER: DX0221KQIGC
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 908-298-2987
; TELEFAX: 908-298-5388
; INFORMATION FOR SEQ ID NO: 42:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 33 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (oligonucleotide)
; US-08-481-560-42

; APPLICATION NUMBER:
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Jane Massey Licata
; REGISTRATION NUMBER: 32,257
; REFERENCE/DOCKET NUMBER: TCEL-00039
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (609) 779-2400
; TELEFAX: (609) 779-8488
; INFORMATION FOR SEQ ID NO: 9:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 31
; TYPE: Nucleic Acid
; STRANDEDNESS: Single
; TOPOLOGY: Linear
; ANTI-SENSE: NO
; PCT-US96-08142-9

Query Match 1.9%; Score 31; DB 5; Length 31;
Best Local Similarity 100.0%; Pred. No. 2.8e+02; Indels 0; Gaps 0;
Matches 31; Conservative 0; Mismatches 0;

QY 119 TGGCAGCCTTCCTCATTTCTGCAGCTCTGTG 149
Db 31 TGGCAGCCTTCCTCATTTCTGCAGCTCTGTG 1

RESULT 9
US-09-308-160B-6/c
; Sequence 6, Application US/09308160B
; Patent No. 6355775
; GENERAL INFORMATION:
; APPLICANT: Nagasawa, Yasuo
; APPLICANT: Yoshida, Hideaki
; TITLE OF INVENTION: TRANSCRIPTIONAL INHIBITOR
; FILE REFERENCE: 4001-0002
; CURRENT APPLICATION NUMBER: US/09/308,160B
; CURRENT FILING DATE: 1999-06-19
; PRIOR APPLICATION NUMBER: PCT/JP97/04127
; PRIOR FILING DATE: 1997-11-12
; PRIOR APPLICATION NUMBER: 8-305043
; PRIOR FILING DATE: 1996-11-15
; NUMBER OF SEQ ID NOS: 31
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 6
; LENGTH: 30
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Primer
; US-09-308-160B-6

Query Match 1.8%; Score 30; DB 4; Length 30;
Best Local Similarity 100.0%; Pred. No. 4.5e+02; Indels 0; Gaps 0;
Matches 30; Conservative 0; Mismatches 0;

QY 14 GACAGCAGCAGCACACAGCTTCTAGGACAA 43
Db 30 GACAGCAGCAGCACACAGCTTCTAGGACAA 1

RESULT 10
US-09-456-399-6/c
; Sequence 6, Application US/09456399
; Patent No. 644801
; GENERAL INFORMATION:
; APPLICANT: Institute of Cytosignal Research, Inc.
; TITLE OF INVENTION: Transcriptional Inhibitor
; FILE REFERENCE: SI-802PCT-US
; CURRENT APPLICATION NUMBER: US/09/456,399
; CURRENT FILING DATE: 1999-12-08
; PRIOR APPLICATION NUMBER: JP 1996-305043
; PRIOR FILING DATE: 1996-11-15
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;; PRIOR APPLICATION NUMBER: PCT/JP97/04127  
;; PRIOR FILING DATE: 1997-11-12  
;; PRIOR APPLICATION NUMBER: US 09/308,160  
;; PRIOR FILING DATE: 1999-05-14  
;; NUMBER OF SEQ ID NOS: 15  
;; SOFTWARE: PatentIn Ver. 2.0  
;; SEQ ID NO 6  
;; LENGTH: 30  
;; TYPE: DNA  
;; ORGANISM: Artificial Sequence  
;; FEATURE:  
;; OTHER INFORMATION: Description of Artificial Sequence: an artificially  
US-09-456-399-6  
;; OTHER INFORMATION: synthesized primer sequence

Query Match 1.8%; Score 30; DB 4; Length 30;  
Best Local Similarity 100.0%; Pred. No. 4.5e+02;  
Matches 30; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 14 GACAGCAGACACACAGCTTCTAGGACAA 43  
Db 30 GACAGCAGACACACAGCTTCTAGGACAA 1

RESULT 11  
US-08-859-998-479  
;; Sequence 479, Application US/08859998  
;; Patent No. 5994076  
;; GENERAL INFORMATION:  
;; APPLICANT: Chenchik, Alex  
;; APPLICANT: Jokhadze, George  
;; APPLICANT: Bibilashvilli, Robert  
;; TITLE OF INVENTION: METHOD OF ASSAYING DIFFERENTIAL  
;; TITLE OF INVENTION: EXPRESSION  
;; NUMBER OF SEQUENCES: 1375  
;; CORRESPONDENCE ADDRESS:  
;; ADDRESSEE: Fish & Richardson, P.C.  
;; STREET: 2200 Sand Hill Road, Suite 100  
;; CITY: Menlo Park  
;; STATE: CA  
;; COUNTRY: US  
;; ZIP: 94025  
;; COMPUTER READABLE FORM:  
;; MEDIUM TYPE: Diskette  
;; COMPUTER: IBM Compatible  
;; OPERATING SYSTEM: Windows95  
;; SOFTWARE: FastSeq for Windows Version 2.0  
;; CURRENT APPLICATION DATA:  
;; APPLICATION NUMBER: US/08/859,998  
;; FILING DATE: 21-MAY-1997  
;; CLASSIFICATION: 435  
;; PRIOR APPLICATION DATA:  
;; APPLICATION NUMBER:  
;; FILING DATE:  
;; ATTORNEY/AGENT INFORMATION:  
;; NAME: Field, Bret E.  
;; REGISTRATION NUMBER: 37,620  
;; REFERENCE/DOCKET NUMBER: 09096/002001  
;; TELECOMMUNICATION INFORMATION:  
;; TELEPHONE: 415-322-5070  
;; TELEFAX: 415-854-0875  
;; INFORMATION FOR SEQ ID NO: 479:  
;; SEQUENCE CHARACTERISTICS:  
;; LENGTH: 28 base pairs  
;; TYPE: nucleic acid  
;; STRANDEDNESS: single  
;; TOPOLOGY: linear  
;; MOLECULE TYPE: DNA  
;; FEATURE:  
;; OTHER INFORMATION: oligonucleotide primer  
US-08-859-998-479

Query Match 1.7%; Score 28; DB 2; Length 28;

Best Local Similarity 100.0%; Pred. No. 1.1e+03;  
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 88 AACATGACTTCCAAAGTCGGCGTGGCTC 115  
Db 1 AACATGACTTCCAAAGTCGGCGTGGCTC 28

RESULT 12  
US-08-859-998-480/c  
;; Sequence 480, Application US/08859998  
;; Patent No. 5994076  
;; GENERAL INFORMATION:  
;; APPLICANT: Chenchik, Alex  
;; APPLICANT: Jokhadze, George  
;; APPLICANT: Bibilashvilli, Robert  
;; TITLE OF INVENTION: METHOD OF ASSAYING DIFFERENTIAL  
;; TITLE OF INVENTION: EXPRESSION  
;; NUMBER OF SEQUENCES: 1375  
;; CORRESPONDENCE ADDRESS:  
;; ADDRESSEE: Fish & Richardson, P.C.  
;; STREET: 2200 Sand Hill Road, Suite 100  
;; CITY: Menlo Park  
;; STATE: CA  
;; COUNTRY: US  
;; ZIP: 94025  
;; COMPUTER READABLE FORM:  
;; MEDIUM TYPE: Diskette  
;; COMPUTER: IBM Compatible  
;; OPERATING SYSTEM: Windows95  
;; SOFTWARE: FastSeq for Windows Version 2.0  
;; CURRENT APPLICATION DATA:  
;; APPLICATION NUMBER: US/08/859,998  
;; FILING DATE: 21-MAY-1997  
;; CLASSIFICATION: 435  
;; PRIOR APPLICATION DATA:  
;; APPLICATION NUMBER:  
;; FILING DATE:  
;; ATTORNEY/AGENT INFORMATION:  
;; NAME: Field, Bret E.  
;; REGISTRATION NUMBER: 37,620  
;; REFERENCE/DOCKET NUMBER: 09096/002001  
;; TELECOMMUNICATION INFORMATION:  
;; TELEPHONE: 415-322-5070  
;; TELEFAX: 415-854-0875  
;; INFORMATION FOR SEQ ID NO: 480:  
;; SEQUENCE CHARACTERISTICS:  
;; LENGTH: 28 base pairs  
;; TYPE: nucleic acid  
;; STRANDEDNESS: single  
;; TOPOLOGY: linear  
;; MOLECULE TYPE: DNA  
;; FEATURE:  
;; OTHER INFORMATION: oligonucleotide primer  
US-08-859-998-480

Query Match 1.7%; Score 28; DB 2; Length 28;  
Best Local Similarity 100.0%; Pred. No. 1.1e+03;  
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 248 GAGTGATTGAGAGTGACACACTGCGC 275  
Db 28 GAGTGATTGAGAGTGACACACTGCGC 1

RESULT 13  
US-09-225-928-479  
;; Sequence 479, Application US/09225928  
;; Patent No. 6352829  
;; GENERAL INFORMATION:  
;; APPLICANT: Chenchik, Alex  
;; APPLICANT: Jokhadze, George  
;; APPLICANT: Bibilashvilli, Robert

SOFTWARE: FASTSEQ FOR WINDOWS VERSION 2.0

LENGTH: 25 base pairs  
TYPE: nucleic acid

COMPUTER: IBM Compatible  
OPERATING SYSTEM: Windows95

COMPUTER: IBM Compatible  
OPERATING SYSTEM: Windows95  
VERSION: 3.0



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; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
US-08-327-494A-8

Query Match      1.5%; Score 25; DB 1; Length 25;
Best Local Similarity 100.0%; Pred. No. 4.6e+03;
Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 358 GAGAGAGTTTGAAGAGGGCTGAGA 382
Db 25 GAGAGAGTTTGAAGAGGGCTGAGA 1

RESULT 16
PCT-US95-13659-8/c
Sequence 8 Application PC/TUS9513659
GENERAL INFORMATION:
APPLICANT: Blaser, Martin J.
APPLICANT: Tumuru, Murali K.R.
APPLICANT: Sharma, Smita A
TITLE OF INVENTION: cagB and cagC Genes for H. pylori and
TITLE OF INVENTION: Related Methods and Compositions
NUMBER OF SEQUENCES: 8
CORRESPONDENCE ADDRESS:
ADDRESSEE: NEEDLE & ROSENBERG, P.C.
STREET: 127 Peachtree Street, Suite 1200
CITY: Atlanta
STATE: Georgia
COUNTRY: USA
ZIP: 303-3
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: PCT/US95/13659
FILING DATE:
CLASSIFICATION:
ATTORNEY/AGENT INFORMATION:
NAME: Spratt, Gwendolyn D.
REGISTRATION NUMBER: 36,016
REFERENCE/DOCKET NUMBER: 2200.029
TELECOMMUNICATION INFORMATION:
TELEPHONE: 404/688-0770
TELEFAX: 404/688-9880
INFORMATION FOR SEQ ID NO: 8:
SEQUENCE CHARACTERISTICS:
LENGTH: 25 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
PCT-US95-13659-8

Query Match      1.5%; Score 25; DB 5; Length 25;
Best Local Similarity 100.0%; Pred. No. 4.6e+03;
Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 358 GAGAGAGTTTGAAGAGGGCTGAGA 382
Db 25 GAGAGAGTTTGAAGAGGGCTGAGA 1

RESULT 17
PCT-US96-08142-8/c
Sequence 8 Application PC/TUS9608142
GENERAL INFORMATION:
APPLICANT: Constance Emmett, Kimberly A. Foster
TITLE OF INVENTION: Universal Chemistry Enzyme-Linked
TITLE OF INVENTION: Immunosorbent Assay for Detection of
TITLE OF INVENTION: mRNA Expression
```

```
; NUMBER OF SEQUENCES: 9
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Jane Massey Licata, Esq.
; STREET: 210 Lake Drive East, Suite 201
; CITY: Cherry Hill
; STATE: NJ
; COUNTRY: USA
; ZIP: 08002
; COMPUTER READABLE FORM:
; MEDIUM TYPE: DISKETTE, 3.5 INCH, 1.44 Mb STORAGE
; COMPUTER: IBM 486
; OPERATING SYSTEM: WINDOWS FOR WORKGROUPS
; SOFTWARE: WORDPERFECT 5.1
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: PCT/US96/08142
; FILING DATE: Herewith
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER:
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Jane Massey Licata
; REGISTRATION NUMBER: 32,257
; REFERENCE/DOCKET NUMBER: TCEL-0039
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (609) 779-2400
; TELEFAX: (609) 779-8488
; INFORMATION FOR SEQ ID NO: 8:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 25
; TYPE: Nucleic Acid
; STRANDEDNESS: Single
; TOPOLOGY: Linear
; ANTI-SENSE: NO
PCT-US96-08142-8

Query Match      1.5%; Score 25; DB 5; Length 25;
Best Local Similarity 100.0%; Pred. No. 4.6e+03;
Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 358 GAGAGAGTTTGAAGAGGGCTGAGA 382
Db 25 GAGAGAGTTTGAAGAGGGCTGAGA 1

RESULT 18
US-09-710-200-72/c
Sequence 72, Application US/09710200
GENERAL INFORMATION:
APPLICANT: Nanogen, Inc.
APPLICANT: Weidenhammer, Elaine M.
APPLICANT: Wang, Ling
APPLICANT: Xu, Xiao
APPLICANT: Heller, Michael J.
APPLICANT: Kahl, Brenda F.
TITLE OF INVENTION: IMPROVED METHODS FOR GENE EXPRESSION MONITORING ON ELECTRONIC
FILE REFERENCE: 256/262 Patrick S. Eagelman
CURRENT APPLICATION NUMBER: US/09/710,200
CURRENT FILING DATE: 2000-11-09
NUMBER OF SEQ ID NOS: 73
SOFTWARE: PatentIn version 3.1
SEQ ID NO 72
LENGTH: 24
TYPE: DNA
ORGANISM: Homo sapiens
US-09-710-200-72

Query Match      1.5%; Score 24; DB 4; Length 24;
Best Local Similarity 100.0%; Pred. No. 7.2e+03;
Matches 24; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 475 GTCTGGGCTCTGTTGTAGGTTGCC 498
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Thu Feb 13 18:20:42 2003

```
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Db 24 GTGTGGCTGCTGTTAGGGTTGCC 1

RESULT 19
PCT-US96-08142-7
; Sequence 7, Application PC/TUS9608142
; GENERAL INFORMATION:
; APPLICANT: Constance Emmett, Kimberly A. Foster
; TITLE OF INVENTION: Universal Chemistry Enzyme-Linked
; TITLE OF INVENTION: Immunosorbent Assay for Detection of
; TITLE OF INVENTION: mRNA Expression
; NUMBER OF SEQUENCES: 9
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Jane Massey Licata, Esq.
; STREET: 210 Lake Drive East, Suite 201
; CITY: Cherry Hill
; STATE: NJ
; COUNTRY: USA
; ZIP: 08002
; COMPUTER READABLE FORM:
; MEDIUM TYPE: DISKETTE, 3.5 INCH, 1.44 MB STORAGE
; COMPUTER: IBM 486
; OPERATING SYSTEM: WINDOWS FOR WORKGROUPS
; SOFTWARE: WORDPERFECT 5.1
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: PCT/US96/08142
; FILING DATE: Herewith
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER:
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Jane Massey Licata
; REGISTRATION NUMBER: 32,257
; REFERENCE/DOCKET NUMBER: TCEL-0039
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (609) 779-2400
; TELEFAX: (609) 779-8488
; INFORMATION FOR SEQ ID NO: 7:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 24
; TYPE: Nucleic Acid
; STRANDEDNESS: Single
; TOPOLOGY: Linear
; ANTI-SENSE: No
PCT-US96-08142-7
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Query Match 1.5%; Score 24; DB 5; Length 24;
Best Local Similarity 100.0%; Pred. No. 7.2e+03;
Matches 24; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 91 ATGACTTCCAAGCTGGCGCTGGCT 114
Db 1 ATGACTTCCAAGCTGGCGCTGGCT 24

RESULT 20
US-09-641-638-1167/c
; Sequence 1167, Application US/09641638
; Patent No. 6432648
; GENERAL INFORMATION:
; APPLICANT: Blumenfeld, Marta
; APPLICANT: Bougueleret, Lydie
; APPLICANT: Chumakov, Ilya
; APPLICANT: Cohen, Annick
; TITLE OF INVENTION: BIALLELIC MARKERS DERIVED FROM GENOMIC REGIONS CARRYING
; TITLE OF INVENTION: GENES INVOLVED IN ARACHIDONIC ACID METABOLISM
; FILE REFERENCE: GENSET.051CPI
; CURRENT APPLICATION NUMBER: US/09/641,638
; CURRENT FILING DATE: 2000-08-16
; PRIOR APPLICATION NUMBER: US 09/502,330
; PRIOR FILING DATE: 2000-02-11
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GenCore version 5.1.3  
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OM nucleic - nucleic search, using sw model

Run on: February 9, 2003, 17:25:52 : Search time 133 seconds  
(without alignments)  
5841.285 Million cell updates/sec

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Perfect score: 1639  
Sequence: 1 acaacttccagacagca.....ataaattgttgcagaagt 1639

Scoring table: IDENTITY\_NUC  
Gapop 10.0 , Gapext 1.0

Searched: 408267 seqs, 237001491 residues

Total number of hits satisfying chosen parameters: 231544

Minimum DB seq length: 0  
Maximum DB seq length: 50

Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
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C 2	30	1.8	30	10	US-09-816-124-4
C 3	25	1.5	25	10	US-09-885-441-31
C 4	24	1.5	24	10	US-09-975-408-72
C 5	24	1.5	24	12	US-10-075-579-72
C 6	21.8	1.3	47	9	US-09-853-526-250
C 7	21.8	1.3	47	9	US-09-853-526-327
C 8	21.8	1.3	47	10	US-09-901-484A-250
C 9	21.8	1.3	47	10	US-09-901-484A-327
C 10	21.6	1.3	41	10	US-09-955-462A-1
C 11	20.4	1.2	34	10	US-09-955-462A-2
C 12	20.4	1.2	48	9	US-09-864-785-3472
C 13	20.2	1.2	48	9	US-09-864-785-3568
C 14	20	1.2	20	10	US-09-975-408-73
C 15	20	1.2	20	12	US-10-075-579-73
C 16	20	1.2	28	10	US-09-142-593-40
C 17	20	1.2	48	9	US-09-864-785-3460
C 18	19.8	1.2	43	9	US-09-828-523A-96
C 19	19.8	1.2	48	9	US-09-825-805-64

C	20	19.6	1.2	45	9	US-10-017-736-9	Sequence 9, Appli
	21	19.4	1.2	49	10	US-09-773-385-3	Sequence 3, Appli
	22	19.4	1.2	50	9	US-09-992-598-45	Sequence 45, Appl
	23	19.4	1.2	50	9	US-09-989-293A-45	Sequence 45, Appl
	24	19.4	1.2	50	9	US-09-989-735-45	Sequence 45, Appl
	25	19.4	1.2	50	9	US-09-990-444-45	Sequence 45, Appl
	26	19.4	1.2	50	9	US-09-989-730-45	Sequence 45, Appl
	27	19.4	1.2	50	9	US-09-990-436-45	Sequence 45, Appl
	28	19.4	1.2	50	9	US-09-991-181-45	Sequence 45, Appl
	29	19.4	1.2	50	9	US-09-993-687-45	Sequence 45, Appl
	30	19.4	1.2	50	9	US-09-989-734-45	Sequence 45, Appl
	31	19.4	1.2	50	9	US-09-997-653-45	Sequence 45, Appl
	32	19.4	1.2	50	9	US-09-993-667-45	Sequence 45, Appl
	33	19.4	1.2	50	10	US-09-989-722-45	Sequence 45, Appl
	34	19.4	1.2	50	10	US-09-989-723-45	Sequence 45, Appl
	35	19.4	1.2	50	10	US-09-989-279-45	Sequence 45, Appl
	36	19.4	1.2	50	10	US-09-989-727-45	Sequence 45, Appl
	37	19.4	1.2	50	10	US-09-989-731-45	Sequence 45, Appl
	38	19.4	1.2	50	10	US-09-989-732-45	Sequence 45, Appl
	39	19.4	1.2	50	10	US-09-991-073-45	Sequence 45, Appl
	40	19.4	1.2	50	10	US-09-990-442-45	Sequence 45, Appl
	41	19.4	1.2	50	10	US-09-991-163-45	Sequence 45, Appl
	42	19.4	1.2	50	10	US-09-993-604-45	Sequence 45, Appl
	43	19.4	1.2	50	10	US-09-990-456-45	Sequence 45, Appl
	44	19.4	1.2	50	10	US-09-989-721-45	Sequence 45, Appl
	45	19.2	1.2	24	10	US-09-885-441-30	Sequence 30, Appl

ALIGNMENTS

RESULT 1  
US-09-801-274-1358  
; Sequence 1358, Application US/09801274  
; Patent No. US2002032319A1  
; GENERAL INFORMATION:  
; APPLICANT: Cargill, Michele  
; APPLICANT: Ireland, James S.  
; APPLICANT: Lander, Eric S.  
; TITLE OF INVENTION: HUMAN SINGLE NUCLEOTIDE POLYMORPHISMS  
; FILE REFERENCE: 2825.2009-001  
; CURRENT APPLICATION NUMBER: US/09/801,274  
; CURRENT FILING DATE: 2001-03-07  
; PRIOR APPLICATION NUMBER: US 60/187,510  
; PRIOR FILING DATE: 2000-03-07  
; PRIOR APPLICATION NUMBER: US 60/206,129  
; PRIOR FILING DATE: 2000-05-22  
; NUMBER OF SEQ ID NOS: 1802  
; SOFTWARE: FastSeq for Windows Version 4.0  
; SEQ ID NO 1358  
; LENGTH: 31  
; TYPE: DNA  
; ORGANISM: Homo sapiens  
US-09-801-274-1358  
  
Query Match 1.9%; Score 30.6; DB 10; Length 31;  
Best Local Similarity 96.8%; Pred. No. 2.4e+03;  
Matches 30; Conservative 1; Mismatches 0; Indels 0; Gaps 0;  
  
QY 220 TTCCACCCCAAAATTTATCAAGAACTGAGAG 250  
Db 1 TTCCACCCCAAAATTTATCAAGAACTGAGAG 31  
  
RESULT 2  
US-09-816-124-4  
; Sequence 4, Application US/09816124  
; Patent No. US20020150897A1  
; GENERAL INFORMATION:  
; APPLICANT: Nagasawa, Yasuo  
; APPLICANT: Yoshida, Hideaki  
; TITLE OF INVENTION: Method for detecting and isolating genes  
; FILE REFERENCE: SI-801PCT

; CURRENT APPLICATION NUMBER: US/09/816.124  
; CURRENT FILING DATE: 2001-03-26  
; PRIOR APPLICATION NUMBER: PCT/JP97/04126  
; PRIOR FILING DATE: 1997-11-12  
; PRIOR APPLICATION NUMBER: JP 1996-305163  
; PRIOR FILING DATE: 1996-11-15  
; NUMBER OF SEQ ID NOS: 4  
; SOFTWARE: PatentIn version 3.0  
; SEQ ID NO 4  
; LENGTH: 30  
; TYPE: DNA  
; ORGANISM: Artificial Sequence  
; FEATURE:  
; OTHER INFORMATION: Artificially synthesized primer sequence  
US-09-816-124-4

Query Match 1.8%; Score 30; DB 10; Length 30;  
Best Local Similarity 100.0%; Pred. No. 3e+03;  
Matches 30; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 14 GACAGCAGCAGCACACAAGCTTCTAGGACAA 43  
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Db 30 GACAGCAGCAGCACACAAGCTTCTAGGACAA 1

RESULT 3  
US-09-885-441-31/c  
; Sequence 11 Application US/09885441  
; Patent No. US20020146407A1  
; GENERAL INFORMATION:  
; APPLICANT: Xiao, Yonghong  
; TITLE OF INVENTION: Regulation of Human Eosinophil Serine  
; FILE REFERENCE: 04974.00512  
; CURRENT APPLICATION NUMBER: US/09/885.441  
; CURRENT FILING DATE: 2001-06-21  
; PRIOR APPLICATION NUMBER: US 60/212.844  
; PRIOR FILING DATE: 2000-06-21  
; PRIOR APPLICATION NUMBER: US 60/244.171  
; PRIOR FILING DATE: 2000-10-31  
; PRIOR APPLICATION NUMBER: US 60/279.766  
; PRIOR FILING DATE: 2001-03-30  
; PRIOR APPLICATION NUMBER: PCT/  
; PRIOR FILING DATE: 2001-06-20  
; NUMBER OF SEQ ID NOS: 58  
; SEQ ID NO 31  
; LENGTH: 25  
; TYPE: DNA  
; ORGANISM: Homo sapiens  
US-09-885-441-31

Query Match 1.5%; Score 25; DB 10; Length 25;  
Best Local Similarity 100.0%; Pred. No. 2.2e+04;  
Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 888 TGTTCACGTGCGCTGGTTCTCC 912  
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Db 25 TGTTCACGTGCGCTGGTTCTCC 1

RESULT 4  
US-09-975-408-72/c  
; Sequence 7 Application US/09975408  
; Patent No. US20020150917A1  
; GENERAL INFORMATION:  
; APPLICANT: Nanogen, Inc.  
; APPLICANT: Weidenhammer, Elaine M.  
; APPLICANT: Xu, Xiao  
; APPLICANT: Kahl, Brenda F.  
; TITLE OF INVENTION: IMPROVED METHODS FOR GENE EXPRESSION MONITORING ON ELECTRONIC MIC  
; FILE REFERENCE: 267/174 Patrick S. Eaglenan  
; CURRENT APPLICATION NUMBER: US/09/975.408

; CURRENT FILING DATE: 2001-10-10  
; PRIOR APPLICATION NUMBER: 09/710.200  
; PRIOR FILING DATE: 2000-11-09  
; NUMBER OF SEQ ID NOS: 73  
; SOFTWARE: PatentIn version 3.1  
; SEQ ID NO 72  
; LENGTH: 24  
; TYPE: DNA  
; ORGANISM: Homo sapiens  
US-09-975-408-72

Query Match 1.5%; Score 24; DB 10; Length 24;  
Best Local Similarity 100.0%; Pred. No. 3.3e+04;  
Matches 24; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 475 GTGTGGGTCTGTGTAGGGTTGCC 498  
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Db 24 GTGTGGGTCTGTGTAGGGTTGCC 1

RESULT 5  
US-10-075-579-72/c  
; Sequence 72 Application US/10075579  
; Patent No. US20020119484A1  
; GENERAL INFORMATION:  
; APPLICANT: Nanogen, Inc.  
; APPLICANT: Weidenhammer, Elaine M.  
; APPLICANT: Wang, Ling  
; APPLICANT: Xu, Xiao  
; APPLICANT: Heller, Michael J.  
; APPLICANT: Kahl, Brenda F.  
; TITLE OF INVENTION: IMPROVED METHODS FOR GENE EXPRESSION MONITORING ON ELECTRONIC  
; FILE REFERENCE: 256/262 Patrick S. Eaglenan  
; CURRENT APPLICATION NUMBER: US/10/075.579  
; CURRENT FILING DATE: 2002-02-12  
; PRIOR APPLICATION NUMBER: US/09/710.200  
; PRIOR FILING DATE: 2000-11-09  
; NUMBER OF SEQ ID NOS: 73  
; SOFTWARE: PatentIn version 3.1  
; SEQ ID NO 72  
; LENGTH: 24  
; TYPE: DNA  
; ORGANISM: Homo sapiens  
US-10-075-579-72

Query Match 1.5%; Score 24; DB 12; Length 24;  
Best Local Similarity 100.0%; Pred. No. 3.3e+04;  
Matches 24; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 475 GTGTGGGTCTGTGTAGGGTTGCC 498  
|||||  
Db 24 GTGTGGGTCTGTGTAGGGTTGCC 1

RESULT 6  
US-09-853-526-250/c  
; Sequence 250 Application US/09853526  
; Patent No. US20020165345A1  
; GENERAL INFORMATION:  
; APPLICANT: Blumenfeld, Marta  
; APPLICANT: Ilya, Chumakov  
; APPLICANT: Bougueret, Lydie  
; TITLE OF INVENTION: PROSTATE CANCER GENE  
; FILE REFERENCE: GENSET 18CPICP  
; CURRENT APPLICATION NUMBER: US/09/853.526  
; CURRENT FILING DATE: 2001-05-11  
; PRIOR APPLICATION NUMBER: 09/338.907  
; PRIOR FILING DATE: 1999-06-23  
; PRIOR APPLICATION NUMBER: 08/996.306  
; PRIOR FILING DATE: 1997-12-22  
; PRIOR APPLICATION NUMBER: 60/099.658  
; PRIOR FILING DATE: 1998-09-09

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RESULT 9
US-09-901-484A-327/c
: Sequence 327, Application US/09901484A
: Patent No. US20020119460A1
: GENERAL INFORMATION:
: APPLICANT: Cohen, Daniel
: APPLICANT: Blumenfeld, Marta
: APPLICANT: Chumakov, Ilya
: APPLICANT: Bouqueleret, Lydie
: TITLE OF INVENTION: Prostate Cancer Gene

```

QY 1038 TATTTATTTATGTAATTTAAGC 1065

Query Match 1.38; Score 21.6; DB 10; Length 41;  
Best Local Similarity 85.7%; Pred. No. 9.6e+04;  
Matches 24; Conservative 0; Mismatches 4; Indels 0; Caps 0;

RESULT 13  
US-09-864-785-3568



## TELECOMMUNICATION INFORMATION:

TELEPHONE: 612-305-1226  
TELEFAX: 612-305-1228  
INFORMATION FOR SEQ ID NO: 40:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 28 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
MOLECULE TYPE: DNA (genomic)  
US-09-142-593-40

Query Match 1.2%; Score 20; DB 10; Length 28;

Best Local Similarity 82.1%; Pred. No. 1.7e+05;  
Matches 23; Conservative 0; Mismatches 5; Indels 0; Gaps 0;

QY 75 CTCACGTGTGTAAACATGACTTCCAAAG 102

Db 28 CTTATTGTATGTAACCTTGACTTTCAG 1

## RESULT 17

US-09-864-785-3460  
Sequence 3460, Application US/09864785  
Patent No. US20020177568A1  
GENERAL INFORMATION:  
APPLICANT: Ribozyme Pharmaceuticals, Inc.  
APPLICANT: Stinchcomb, Dan  
APPLICANT: Draper, Ken  
APPLICANT: McSwiggen, Jim

TITLE OF INVENTION: Enzymatic Nucleic Acid Treatment of Diseases or Conditions Related to NF-Kappa B  
TITLE OF INVENTION: Levels of NF-Kappa B  
FILE REFERENCE: 400/022 (MBH00-812-D)  
CURRENT APPLICATION NUMBER: US/09/864,785  
CURRENT FILING DATE: 2001-05-23  
NUMBER OF SEQ ID NOS: 3929  
SOFTWARE: PatentIn version 3.0  
SEQ ID NO 3460  
LENGTH: 48  
TYPE: RNA  
ORGANISM: Artificial Sequence  
FEATURE:  
OTHER INFORMATION: Description of Artificial Sequence: Nucleic Acid

US-09-864-785-3460

Query Match 1.2%; Score 20; DB 9; Length 48;

Best Local Similarity 61.4%; Pred. No. 1.9e+05;  
Matches 27; Conservative 2; Mismatches 15; Indels 0; Gaps 0;

QY 13 AGACAGCAGCACACAAAGCTTCTAGGACAGAGCCGAGGAAGAA 56

Db 2 AGACUCCGAGGAACUCCUUCAGGACACUCCGCGGACAAA 45

## RESULT 18

US-09-828-523A-96  
Sequence 96, Application US/09828523A  
Patent No. US20020168697A1  
GENERAL INFORMATION:  
APPLICANT: The Pharmacia & Upjohn Company  
TITLE OF INVENTION: ANTIMICROBIAL METHODS AND MATERIALS  
FILE REFERENCE: 268.62120101  
CURRENT APPLICATION NUMBER: US/09/828,523A  
CURRENT FILING DATE: 2001-04-06  
PRIOR APPLICATION NUMBER: 60/266,327  
PRIOR FILING DATE: 2000-04-06  
NUMBER OF SEQ ID NOS: 99  
SOFTWARE: PatentIn version 3.1  
SEQ ID NO 96  
LENGTH: 43  
TYPE: DNA  
ORGANISM: Artificial Sequence  
FEATURE:  
OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid



OTHER INFORMATION: n stands for inverted deoxyabasic derivative  
US-09-825-805-64

Query Match 1.2%; Score 19.8; DB 9; Length 48;  
Best Local Similarity 59.6%; Pred. No. 2e+05;  
Matches 28; Conservative 2; Mismatches 17; Indels 0; Gaps 0;  
Qy 10 CAGACAGACAGACACACAGCTTCTAGGACAGAGCCAGGAGAA 56  
| | | | | | | | | | | | | | | | | | | | | | | | | |  
Db 1 CGGGUUAUCGGAGGAACUCCUUCAGGACAUCCGUCGGGAGAA 47

RESULT 20  
US-10-017-736-9/c  
; Sequence 9, Application US/10017736  
; Publication No. US20020192640A1  
; GENERAL INFORMATION:  
; APPLICANT: Boehringer Ingelheim (Canada) Ltd.  
; TITLE OF INVENTION: Purified Active HCV NS2/3 Protease  
; FILE REFERENCE: 13/082  
; CURRENT APPLICATION NUMBER: US/10/017.736  
; CURRENT FILING DATE: 2001-12-14  
; PRIOR APPLICATION NUMBER: 60/256,031  
; PRIOR FILING DATE: 2000-12-15  
; NUMBER OF SEQ ID NOS: 21  
; SOFTWARE: FastSEQ for Windows Version 4.0  
; SEQ ID NO 9  
; LENGTH: 45  
; TYPE: DNA  
; ORGANISM: HCV  
US-10-017-736-9

Query Match 1.2%; Score 19.6; DB 9; Length 45;  
Best Local Similarity 66.7%; Pred. No. 2.2e+05;  
Matches 28; Conservative 0; Mismatches 14; Indels 0; Gaps 0;  
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| | | | | | | | | | | | | | | | | | | | | | | | | |  
Db 45 ATGGTGATGCTGCGAGCTTTTCTTTTCATATGGCA 4

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Job time : 136 secs

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Copyright (c) 1993 - 2003 Compugen Ltd.

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(without alignments)  
11238.104 Million cell updates/sec

Title: US-09-960-143-3  
Perfect score: 1639  
Sequence: 1 acaaaacttcagacagca.....atataattgtgtcaaaagt 1639

Scoring table: IDENTITY\_NUC  
Gapop 10.0 , Gapext 1.0

Searched: 16154066 seqs, 8097743376 residues

Total number of hits satisfying chosen parameters: 102860

Minimum DB seq length: 0  
Maximum DB seq length: 50

Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

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1: em\_estba: \*  
2: em\_esthum: \*  
3: em\_estin: \*  
4: em\_estnu: \*  
5: em\_estov: \*  
6: em\_estpl: \*  
7: em\_estro: \*  
8: em\_htc: \*  
9: gb\_est1: \*  
10: gb\_est2: \*  
11: gb\_htc: \*  
12: gb\_est3: \*  
13: gb\_est4: \*  
14: gb\_est5: \*  
15: em\_estfun: \*  
16: em\_estom: \*  
17: gb\_gss: \*  
18: em\_gss\_hum: \*  
19: em\_gss\_inv: \*  
20: em\_gss\_pin: \*  
21: em\_gss\_vrt: \*  
22: em\_gss\_fun: \*  
23: em\_gss\_mam: \*  
24: em\_gss\_mus: \*  
25: em\_gss\_other: \*  
26: em\_gss\_pro: \*  
27: em\_gss\_rod: \*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	23.8	1.5	43	17	AZ404728
2	23.2	1.4	46	13	BJ015738
C 3	23	1.4	50	12	BF029726
C 4	22.8	1.4	46	14	BQ704909
5	22.8	1.4	49	13	BJ054665
C 6	22.8	1.4	49	17	AZ333209

7	22.6	1.4	46	9	AU009443
8	22.6	1.4	46	9	AU009444
9	22.4	1.4	48	9	AU266602
10	22.4	1.4	48	17	TA266090
C 11	22.4	1.4	49	17	AZ346760
12	22.4	1.4	50	10	AW333744
13	22.2	1.4	40	17	AZ615013
14	22.2	1.4	48	13	BJ078372
15	22	1.3	50	9	AU268417
16	21.6	1.3	49	12	BG179823
C 17	21.6	1.3	49	13	BI745472
C 18	21.6	1.3	50	10	AV957350
19	21.4	1.3	44	17	BH853222
20	21.4	1.3	45	17	BH850240
21	21.2	1.3	47	17	TA48H05P
22	21.2	1.3	34	13	BJ052720
23	21.2	1.3	43	17	AZ586882
24	21.2	1.3	45	9	AU267061
25	21.2	1.3	45	17	BH789466
26	21.2	1.3	45	17	BH799966
C 27	21.2	1.3	45	17	TA344F02P
C 28	21.2	1.3	50	17	BH861292
C 29	21.2	1.3	50	17	TA187C12P
30	21	1.3	39	9	AU267631
C 31	21	1.3	43	14	T25548
32	21	1.3	47	9	AU265820
33	21	1.3	48	13	BJ083775
34	21	1.3	50	9	AU006647
35	21	1.3	50	17	AZ950287
C 36	20.8	1.3	39	9	AU266450
C 37	20.8	1.3	44	10	AV834222
C 38	20.8	1.3	45	14	D34826
C 39	20.8	1.3	48	9	AU263470
C 40	20.8	1.3	48	12	BG667239
C 41	20.8	1.3	48	14	C21451
C 42	20.8	1.3	49	13	BI863578
C 43	20.6	1.3	45	17	BH792269
C 44	20.6	1.3	49	13	BI863578
45	20.6	1.3	50	13	BM126011

ALIGNMENTS

RESULT 1	AZ404728	AZ404728	43 bp	DNA	linear	GSS 03-OCT-2000
LOCUS	IM0173108F	Mouse 10kb plasmid	UUGCIM	library	Mus musculus	genomic
DEFINITION	clone UUGCIM0173108 F, DNA sequence.					
ACCESSION	AZ404728					
VERSION	AZ404728.1	GI:10528741				
KEYWORDS	GSS.					
SOURCE	house mouse.					
ORGANISM	Mus musculus					
REFERENCE	Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus. 1 (bases 1 to 43)					
AUTHORS	Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C., Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly,M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A. and Wright,D., Weiss,R.					
TITLE	Mouse whole genome scaffolding with paired end reads from 10kb plasmid inserts					
JOURNAL	Unpublished (2000)					
COMMENT	Contact: Robert B. Weiss University of Utah Genome Center Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT 84112, USA Tel: 801 585 5606 Fax: 801 585 7177 Email: ddunn@genetics.utah.edu Insert Length: 10000 Std Error: 0.00					





/db\_xref="taxon:10090"  
 /clone="fUGC1M0062L10"  
 /clone\_lib="Mouse 10kb plasmid UUGC1M library"  
 /sex="Male"  
 /lab\_host="E. Coli strain XL10-Gold, T1-resistant, F-"  
 /note="Vector: PWD42nv; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource (http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of pWD42 (g147321141gb/AF129072.1), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptor vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

BASE COUNT 24 a 0 c 0 g 25 t  
 ORIGIN  
 Query Match 1.4%; Score 22.8; DB 17; Length 49;  
 Best Local Similarity 71.4%; Pred. No. 3.1e+06;  
 Matches 30; Conservative 0; Mismatches 12; Indels 0; Gaps 0;

QY 1139 TAAATTTATTTATTTAGATATTAATGATGTTTATTAGA 1180  
 ||| | ||| ||| || ||||| || || ||||| |  
 Db 43 TAATTATATATATATATATATATATTAATATATATATA 2

RESULT 7  
 AU009443  
 LOCUS  
 DEFINITION AU009443 Schizosaccharomyces pombe late log phase cDNA EST 31-JUL-1998  
 Schizosaccharomyces pombe cDNA clone spc04922, mRNA sequence.  
 ACCESSION AU009443  
 VERSION AU009443.1 GI:3346123  
 KEYWORDS EST.  
 SOURCE fission yeast.  
 ORGANISM Schizosaccharomyces pombe  
 Eukaryota; Fungi; Ascomycota; Schizosaccharomycetes;  
 Schizosaccharomycetales; Schizosaccharomycetaceae;  
 Schizosaccharomycetes.  
 1 (bases 1 to 46)  
 Morimyo, M. and Mita, K.  
 Identification of expressed sequence tags of Schizosaccharomyces pombe

REFERENCE 1  
 AUTHORS Morimyo, M. and Mita, K.  
 TITLE Identification of expressed sequence tags of Schizosaccharomyces pombe  
 JOURNAL Unpublished (1998)  
 COMMENT Contact: Mitsuoki Morimyo  
 Genome Research Group  
 National Institute of Radiological Sciences  
 9-1, Anagawa-4-chome, Inage-ku, Chiba, Chiba 263-8555, Japan  
 Email: morimyo@nirs.go.jp.

FEATURES  
 source  
 1..46  
 /organism="Schizosaccharomyces pombe"  
 /strain="972"  
 /db\_xref="taxon:4896"  
 /clone="spc04922"  
 /clone\_lib="Schizosaccharomyces pombe late log phase cDNA"  
 /sex="h minus"  
 /note="Vector: M13mp19; The cDNA library of Schizosaccharomyces pombe was prepared by cloning cDNA into the SmaI site of M13mp19 DNA and the direction of DNA sequences was not always from 5' to 3'. The cDNA data of Schizosaccharomyces pombe are available for searching on the World Wide Web. (URL, http://www.nirs.go.jp)"

BASE COUNT 27 a 2 c 3 g 14 t  
 ORIGIN

Query Match 1.4%; Score 22.6; DB 9; Length 46;  
 Best Local Similarity 68.9%; Pred. No. 3.4e+06;  
 Matches 31; Conservative 0; Mismatches 14; Indels 0; Gaps 0;

QY 582 ATACTTATATGTAAGACTATATTATTTTGAATCTACAAAAACAA 626  
 | | ||||| ||| | | ||||| ||||| ||||| ||  
 Db 2 AAAATTATCTTAAATAATTAATTAATTTGAATTTGACAAAAAANA 46

RESULT 8  
 AU009444  
 LOCUS  
 DEFINITION AU009444 Schizosaccharomyces pombe late log phase cDNA EST 31-JUL-1998  
 Schizosaccharomyces pombe cDNA clone spc04923, mRNA sequence.  
 ACCESSION AU009444  
 VERSION AU009444.1 GI:3346124  
 KEYWORDS EST.  
 SOURCE fission yeast.  
 ORGANISM Schizosaccharomyces pombe  
 Eukaryota; Fungi; Ascomycota; Schizosaccharomycetes;  
 Schizosaccharomycetales; Schizosaccharomycetaceae;  
 Schizosaccharomycetes.  
 1 (bases 1 to 46)  
 Morimyo, M. and Mita, K.  
 Identification of expressed sequence tags of Schizosaccharomyces pombe

REFERENCE 1  
 AUTHORS Morimyo, M. and Mita, K.  
 TITLE Identification of expressed sequence tags of Schizosaccharomyces pombe  
 JOURNAL Unpublished (1998)  
 COMMENT Contact: Mitsuoki Morimyo  
 Genome Research Group  
 National Institute of Radiological Sciences  
 9-1, Anagawa-4-chome, Inage-ku, Chiba, Chiba 263-8555, Japan  
 Email: morimyo@nirs.go.jp.

FEATURES  
 source  
 1..46  
 /organism="Schizosaccharomyces pombe"  
 /strain="972"  
 /db\_xref="taxon:4896"  
 /clone="spc04923"  
 /clone\_lib="Schizosaccharomyces pombe late log phase cDNA"  
 /sex="h minus"  
 /note="Vector: M13mp19; The cDNA library of Schizosaccharomyces pombe was prepared by cloning cDNA into the SmaI site of M13mp19 DNA and the direction of DNA sequences was not always from 5' to 3'. The cDNA data of Schizosaccharomyces pombe are available for searching on the World Wide Web. (URL, http://www.nirs.go.jp)"

BASE COUNT 27 a 2 c 3 g 14 t  
 ORIGIN  
 Query Match 1.4%; Score 22.6; DB 9; Length 46;  
 Best Local Similarity 68.9%; Pred. No. 3.4e+06;  
 Matches 31; Conservative 0; Mismatches 14; Indels 0; Gaps 0;

QY 582 ATACTTATATGTAAGACTATATTATTTTGAATCTACAAAAACAA 626  
 | | ||||| ||| | | ||||| ||||| ||||| ||  
 Db 2 AAAATTATCTTAAATAATTAATTAATTTGAATTTGACAAAAAANA 46

RESULT 9  
 AU266602  
 LOCUS  
 DEFINITION AU266602 VS Dictyostellium discoideum cDNA clone VSG534 5', mRNA EST 10-MAY-2002  
 Schizosaccharomyces pombe cDNA clone VSG534 5', mRNA sequence.  
 ACCESSION AU266602  
 VERSION AU266602.1 GI:20525400  
 KEYWORDS EST.  
 SOURCE Dictyostellium discoideum.  
 ORGANISM Dictyostellium discoideum  
 Eukaryota; Mycetozoa; Dictyostellida; Dictyostellium.  
 1 (bases 1 to 48)







BGI/9823	BGL79823	49 bp	linear	EST 06-FEB-2001
LOCUS	602329028F1	NIH_MGC_91	Homo sapiens	cdna clone IMAGE:4430262 5',
DEFINITION				

JOURNAL  
UNPUBLISHED (1999)  
COMMENT  
Contact: Robert Strausberg, Ph.D.

FEATURES  
Location/Qualifiers  
1. 49  
source

BASE COUNT 32 a 4 c 4 q 9 t

Query Match 1.3%; Score 21.6; DB 12; Length 49;

Best Local Similarity 68.2%; Pred. NO. 5e+06;  
Matches 30; Conservative 0; Mismatches 14; Indels 0; Gaps 0;

db

5 TATATAAATAAAAAACAGCGCTTCCAAATAAAAAAAAAAAAAAAAAA 48

RESULT 17

**DEFINITION** rk81a10.y3 Meloidogynae javanica egg pAMPl v6 Chiapelli McCarter Meloidogynae javanica cDNA 5', mRNA sequence.

KEYWORDS EST.  
SOURCE root -knot nematode

Eukaryota; Metazoa; Nematoda; Chromadorea; Tylenchida; Tylenchina;



QY 1122 TTATAAGATGTTATAGTAAATTTATTTATTTTAGATA 1160  
 || ||||| ||| ||||| ||||| |||||  
 Db 3 TTTAAAGATGACTTAGACATTTTATTTCTTTAAGAAA 41

RESULT 20

BH850240

LOCUS

DEFINITION

BH850240 45 bp DNA linear GSS 13-JUN-2002  
 SALK\_071000.50.25.x Arabidopsis thaliana TDNA insertion lines  
 Arabidopsis thaliana genomic clone SALK\_071000.50.25.x, DNA  
 sequence.

ACCESSION

BH850240

VERSION

BH850240.1

KEYWORDS

GSS.

SOURCE

ORGANISM

Arabidopsis thaliana  
 Arabidopsis thaliana  
 Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;  
 Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;  
 Rosidae; eurosids II; Brassicales; Brassicaceae; Arabidopsis.  
 1 (bases 1 to 45)  
 Alonso,J.M., Leishe,T.J., Barajas,P., Chen,H., Cheuk,R., Gadrinab  
 ,C., Jeske,A., Karnes,M., Kim,C.J., Parker,H., Prednis,L., Shinn,P.  
 , Zimmerman,J. and Ecker,J.R.  
 A Sequence-Indexed Library of Insertion Mutations in the  
 Arabidopsis Genome  
 Unpublished (2001)  
 Contact: Joseph R. Ecker  
 Salk Institute Genomic Analysis Laboratory (SIGnAL)  
 The Salk Institute for Biological Studies  
 10010 N. Torrey Pines Road, La Jolla, CA 92037, USA  
 Tel: 858 453 4100 x1752  
 Fax: 858 558 6379  
 Email: ecker@salk.edu  
 This is single pass sequence recovered from the left border of  
 TDNA.

TITLE

JOURNAL

COMMENT

Class: TDNA tagged.

Location/Qualifiers

1..45

/organism="Arabidopsis thaliana"

/strain="Columbia 0"

/db\_xref="taxon:3702"

/clone="SALK\_071000.50.25.x"

/note="PCR was performed on Arabidopsis thaliana lines  
 each of which contains one or more TDNA insertion  
 elements. The resultant fragment for each line was  
 directly sequenced to determine the genomic sequence at  
 the site of insertion. Details of the protocols used can  
 be found at [http://signal.salk.edu/tdna\\_protocols.html](http://signal.salk.edu/tdna_protocols.html)"

BASE COUNT 16 a 2 c 5 g 22 t

ORIGIN

Query Match 1.3%; Score 21.4; DB 17; Length 45;

Best Local Similarity 71.8%; Pred. No. 5.6e+06;

Matches 28; Conservative 0; Mismatches 11; Indels 0; Gaps 0;

QY 1262 TAATTTTATAGTAAAGTACATTTATTTATCTGAAAT 1300

||||| ||| ||||| ||||| |||||

Db 7 TAATTTTAAAAATAGTGTGATTAATTTGATCTGAATT 45

Search completed: February 9, 2003, 19:20:11  
 Job time : 2368 secs

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GenCore version 5.1.3  
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: February 9, 2003, 15:17:21 : Search time 340 seconds  
(without alignments)  
10855.957 Million cell updates/sec

Title: US-09-960-143-3

Perfect score: 1639

Sequence: 1 acaaaccttcagacagca.....atataattgtgtcagaagt 1639

Scoring table: IDENTITY\_NUC

Gapop 10.0 , Gapext 1.0

Searched: 2185239 seqs, 112599159 residues

Total number of hits satisfying chosen parameters: 2166140

Minimum DB seq length: 0

Maximum DB seq length: 50

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

- N\_Geneseq\_101002:\*
- 1: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1980.DAT:\*
  - 2: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1981.DAT:\*
  - 3: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1982.DAT:\*
  - 4: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1983.DAT:\*
  - 5: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1984.DAT:\*
  - 6: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1985.DAT:\*
  - 7: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1986.DAT:\*
  - 8: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1987.DAT:\*
  - 9: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1988.DAT:\*
  - 10: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1989.DAT:\*
  - 11: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1990.DAT:\*
  - 12: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1991.DAT:\*
  - 13: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1992.DAT:\*
  - 14: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1993.DAT:\*
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  - 21: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA2000.DAT:\*
  - 22: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA2001A.DAT:\*
  - 23: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA2001B.DAT:\*
  - 24: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA2002.DAT:\*

pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match %	Length	ID	Description
1	37	2.3	37	18 AAT76272	Human MDNCF antise
2	37	2.3	37	20 AAX54069	Monocyte-derived n
3	37	2.3	37	21 AAF19635	Human monocyte der
4	37	2.3	37	21 AAA33513	Low adenosine anti
5	34	2.1	34	18 AAT76366	Human interleukin
6	34	2.1	34	20 AAX54167	Human IL-8 antisen
7	34	2.1	34	21 AAF19733	Human interleukin
8	34	2.1	34	21 AAA33611	Low adenosine anti
9	33	2.0	33	14 AAQ37164	Anti-sense primer

C 10	33	2.0	33	17 AAT38979	Interleukin IL-8 3
C 11	33	2.0	33	20 AAV65360	Interleukin-8 anti
C 12	33	2.0	33	20 AAV08015	Primer IL-8 for In
C 13	31	1.9	31	18 AAT45913	Human interleukin-
C 14	31	1.9	31	22 AAI30870	Human single nucle
C 15	30	1.8	30	17 AAT45757	Human interleukin-
C 16	30	1.8	30	19 AAV61828	PCR primer for hum
C 17	30	1.8	30	19 AAV35562	Human interleukin
C 18	30	1.8	30	19 AAV35781	PCR primer IL8p2 o
C 19	30	1.8	30	22 AAC88174	Human interleukin
C 20	30	1.8	30	24 ABL46338	Human interleukin-
C 21	29	1.8	30	19 AAV24558	IL-8 antisense oli
C 22	28	1.7	28	24 ABK66391	Human gene specifi
C 23	28	1.7	28	24 ABK66392	Human gene specifi
C 24	26	1.6	26	24 AAD21908	PCR primer, 2767T
C 25	25.2	1.5	39	18 AAT61103	Chimeric chemokine
C 26	25.2	1.5	39	18 AAT61104	Chimeric chemokine
C 27	25	1.5	25	17 AAT29243	Human interleukin-
C 28	25	1.5	25	18 AAT45912	Human interleukin-
C 29	25	1.5	25	21 AAA65326	B-thromboglobulin
C 30	25	1.5	25	24 AAD39024	Human beta-actin s
C 31	25	1.5	25	24 AAD37345	Interleukin-8 (IL-
C 32	25	1.5	25	24 ABK14346	Human interleukin-
C 33	25	1.5	26	24 AAD21907	Chimeric chemokine
C 34	24.4	1.5	48	18 AAT61099	PCR primer, 2767A
C 35	24	1.5	24	18 AAT45911	Human interleukin-
C 36	24	1.5	24	24 AAD39023	Human beta-actin s
C 37	24	1.5	24	24 AAD37344	Interleukin-8 (IL-
C 38	24	1.5	48	22 AAH30030	Human interleukin
C 39	23.6	1.4	39	18 AAT61109	Chimeric chemokine
C 40	23.6	1.4	39	18 AAT61110	Chimeric chemokine
C 41	23	1.4	23	17 AAT29242	Human interleukin-
C 42	23	1.4	23	18 AAT76365	Human interleukin
C 43	23	1.4	23	20 AAX54166	Human IL-8 antisen
C 44	23	1.4	23	21 AAF19732	Human interleukin-
C 45	23	1.4	23	21 AAA33610	Low adenosine anti

ALIGNMENTS

RESULT 1  
AAT76272/C  
ID AAT76272 standard; DNA; 37 BP.  
XX  
XX  
XX AAT76272;

DT 15-SEP-1997 (first entry)  
XX Human MDNCF antisense oligonucleotide HSMDCNFAS2.  
DE  
XX  
XX Asthma; airway epithelium; adenosine free; cystic fibrosis;  
KW chronic obstructive pulmonary disease; bronchitis;  
KW monocyte-derived neutrophil chemotactic factor; ss.  
XX  
OS Synthetic.  
XX  
XX WO9640162-A1.  
XX  
PD 19-DEC-1996.  
XX  
XX 06-JUN-1996; 96WO-US09306.  
XX  
XX 07-JUN-1995; 95US-0474497.  
PR (UYEC-) UNIV EAST CAROLINA.  
XX  
XX Metzger WJ, Nyce JW;  
XX  
XX WPI; 1997-051871/05.  
XX  
XX Treatment of airway diseases such as asthma - by topically applying  
PT adenosine-free antisense oligo:nucleotide to airway epithelium of

PT subject  
XX Claim 5; Page 33; 71pp; English.  
PS  
XX  
CC A method for treating airway disease in a subject has been produced,  
CC which involves the topical administration of an essentially adenosine  
CC free antisense oligonucleotide (ON) to the airway epithelium of the  
CC subject. The present sequence is an antisense oligonucleotide  
CC HSMNCFAS2 specific for the human monocyte-derived neutrophil  
CC chemotactic factor. The method can be used to treat airway diseases  
CC such as cystic fibrosis, asthma, chronic obstructive pulmonary disease,  
CC bronchitis and other airway diseases characterised by an inflammatory  
CC response. By eliminating adenosine from the antisense ON, its  
CC liberation upon antisense degradation is prevented, thereby preventing  
CC adenosine-induced bronchoconstriction in patients with hyper-reactive  
CC airways.  
XX  
SQ Sequence 37 BP; 0 A; 11 C; 9 G; 17 T; 0 other;  
Query Match 2.3%; Score 37; DB 18; Length 37;  
Best Local Similarity 100.0%; Pred. No. 1.4e+03;  
Matches 37; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 37 AGGACAAGAGCCAGGAGGAAGAACCCGCGAAGGAACCA 73  
Db 37 AGGACAAGAGCCAGGAGGAAGAACCCGCGAAGGAACCA 1  
RESULT 2  
AA54069/c  
ID AAX54069 standard; DNA; 37 BP.  
XX  
AC AAX54069;  
XX  
DT 05-JUL-1999 (first entry)  
XX  
DE Monocyte-derived neutrophil chemotactic factor antisense oligo.  
XX  
KW Antisense oligonucleotide; multiple target; antisense treatment;  
KW impaired respiration; inflammation; lung disease;  
KW pulmonary vasoconstriction; inflammation; allergic rhinitis;  
KW acute asthma; allergy; asthma; impeded respiration;  
KW respiratory distress syndrome; pain; cystic fibrosis;  
KW pulmonary hypertension; pulmonary vasoconstriction; emphysema;  
KW chronic obstructive pulmonary disease; leukemia; lymphoma; carcinoma;  
KW colon cancer; breast cancer; lung cancer; pancreatic cancer;  
KW hepatocellular carcinoma; kidney cancer; melanoma; hepatic metastasis;  
KW prostate cancer; ss.  
XX  
OS Synthetic.  
XX  
PN WO9913886-A1.  
XX  
PD 25-MAR-1999.  
XX  
PF 17-SEP-1998; 98WO-US19419.  
XX  
PR 09-JUN-1998; 98US-0093972.  
XX  
PR 17-SEP-1997; 97US-0059160.  
XX  
PA (UYEC-) UNIV EAST CAROLINA.  
XX  
XX Nyce JW;  
XX  
XX WPI; 1999-229400/19.  
XX  
XX New antisense oligonucleotides used in treatment of, e.g. pulmonary  
XX vasoconstriction  
PT  
XX  
XX Disclosure; Page 51; 120pp; English.  
PS  
XX The specification describes antisense oligonucleotides (AAX52869-X55271)  
XX directed against at least 2 mRNAs selected from target genes, coding and  
CC

CC non-coding regions of RNAs corresponding to target genes, gene  
CC initiation codons, genomic flanking regions, intron-exon borders, the  
CC 5'-end, the 3'-end and the juxta-section between coding and non-coding  
CC regions and all segments of RNAs encoding proteins associated with one  
CC or more diseases, conditions or mixtures. The antisense oligonucleotides  
CC may be derived from sequences AAX55272-74. These multiple target  
CC oligonucleotides (specifically AAX55180-271) can be used for the  
CC antisense treatment of diseases and conditions. Typical diseases and  
CC conditions are those associated with impaired respiration and  
CC inflammation, including lung diseases, pulmonary vasoconstriction,  
CC inflammation, allergic rhinitis, acute asthma, allergies, asthma, impeded  
CC respiration, respiratory distress syndrome, pain, cystic fibrosis,  
CC pulmonary hypertension, pulmonary vasoconstriction, emphysema, chronic  
CC obstructive pulmonary disease (COPD), and cancers such as leukemias,  
CC lymphomas, carcinomas e.g. colon cancer, breast cancer, lung cancer,  
CC pancreatic cancer, hepatocellular carcinoma, kidney cancer, melanoma,  
CC hepatic metastases, as well as all types of cancers which may metastasize  
CC or have metastasized to the lungs, including breast and prostate cancer.  
XX  
SQ Sequence 37 BP; 0 A; 11 C; 9 G; 17 T; 0 other;  
Query Match 2.3%; Score 37; DB 20; Length 37;  
Best Local Similarity 100.0%; Pred. No. 1.4e+03;  
Matches 37; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 37 AGGACAAGAGCCAGGAGGAAGAACCCGCGAAGGAACCA 73  
Db 37 AGGACAAGAGCCAGGAGGAAGAACCCGCGAAGGAACCA 1  
RESULT 3  
AAF19635/c  
ID AAF19635 standard; DNA; 37 BP.  
XX  
AC AAF19635;  
XX  
DT 14-MAR-2001 (first entry)  
XX  
DE Human monocyte derived neutrophil chemotactic factor DNA fragment #1202.  
XX  
KW Low adenosine antisense oligonucleotide; phosphorothioate; allergy;  
KW human; airway disorder; bronchoconstriction; lung inflammation;  
KW surfactant depletion; respiratory; bronchodilator; antiinflammatory;  
KW immunosuppressive; antiasthmatic; analgesic; hypotensive; cytostatic;  
KW respiratory obstruction; pulmonary obstruction; impeded respiration;  
KW surfactant hypoproduction; pulmonary vasoconstriction; impeded respiration;  
KW respiratory distress syndrome; pain; cystic fibrosis; allergic rhinitis;  
KW pulmonary hypertension; emphysema; pulmonary transplantation rejection;  
KW chronic obstructive pulmonary disease; pulmonary infection; bronchitis;  
KW cancer; ss.  
XX  
OS Homo sapiens.  
XX  
PN WO200062736-A2.  
XX  
PD 26-OCT-2000.  
XX  
PF 24-MAR-2000; 2000WO-US08020.  
XX  
PR 06-APR-1999; 99US-0127958.  
XX  
PA (UYEC-) UNIV EAST CAROLINA.  
XX  
PA (NYCE/) NYCE J W.  
XX  
XX Nyce JW;  
XX  
XX WPI; 2000-679539/66.  
XX  
XX Low adenosine (A) content antisense oligonucleotides which do not  
XX trigger adenosine receptors during metabolism, useful e.g. for treating  
XX cancers and respiratory obstructions -  
PS Claim 14; Page 210; 1592pp; English.

XX The present invention describes low adenosine (A) content antisense  
CC oligonucleotides and compositions (I) comprising them. In the antisense  
CC oligonucleotides the A is replaced by a 'universal' or alternative base.  
CC (I) can have respiratory, bronchodilator, antiinflammatory, analgesic,  
CC immunosuppressive, antiasthmatic, hypotensive and cytostatic activities.  
CC The antisense oligonucleotides and (I) can be used to down-regulate the  
CC expression and or activity of target polypeptides associated with  
CC lung/respiratory disorders and malignancies, such as stimulating and  
CC activating peptide factors and transmitters, transcription factors,  
CC immunoglobulins and antibodies, antibody receptors, cytokines and  
CC chemokines, endogenously produced specific and non-specific enzymes,  
CC binding proteins, adhesion molecules and their receptors, cytokine and  
CC chemokine receptors, adenosine receptors, bradykinin receptors, central  
CC nervous system (CNS) and peripheral nervous and non-nervous system  
CC receptors, CNS and peripheral nervous and non-nervous system peptide  
CC transmitters, defensins, growth factors, vasoactive peptides and  
CC receptors, binding proteins and malignancy associated proteins. The  
CC antisense oligonucleotides may be used in this way to treat disorders  
CC including respiratory obstruction (especially pulmonary obstruction  
CC and/or bronchoconstriction) and/or lung inflammation, allergy(ies)  
CC and/or surfactant hypoproduction which are associated with a disease or  
CC condition selected from pulmonary vasoconstriction, inflammation,  
CC allergies, asthma, impeded respiration, respiratory distress syndrome  
CC (RDS), pain, cystic fibrosis (CF), allergic rhinitis (AR), pulmonary  
CC hypertension, emphysema, chronic obstructive pulmonary disease (COPD),  
CC pulmonary transplantation rejection, pulmonary infections, bronchitis,  
CC and/or cancer. AAF18434 to AAF21543 represent human polynucleotide  
CC fragments and antisense oligonucleotides used in the exemplification of  
CC the present invention.

XX Sequence 37 BP; 0 A; 11 C; 9 G; 17 T; 0 other;

Query Match 2.3%; Score 37; DB 21; Length 37;  
Best Local Similarity 100.0%; Pred. No. 1.4e+03;  
Matches 37; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 37 AGGACAAGAGCCAGGAGAAACACCGGAGGAGACCA 73  
Db 37 AGGACAAGAGCCAGGAGAAACACCGGAGGAGACCA 1

RESULT 4  
AAA33513/C  
ID: AAA33513 standard; DNA; 37 BP.

XX AAA33513;

XX 28-JUL-2000 (first entry)

XX Low adenosine antisense oligonucleotide SEQ ID NO:1202.

XX Human: adenosine receptor; low adenosine antisense oligonucleotide;  
KW phosphorothioate; impaired respiration; inflammation; allergy;  
KW allergic disease; bronchoconstriction; inhibitor; antiinflammatory;  
KW antiallergic; antiasthmatic; cytostatic; analgesic; impaired airway;  
KW lung disease; ischaemic condition; pulmonary vasoconstriction; asthma;  
KW respiratory distress syndrome; pain; cystic fibrosis; emphysema;  
KW pulmonary hypertension; chronic obstructive pulmonary disease; COPD;  
KW cancer; leukaemia; lymphoma; carcinoma; metastasis; ss.

XX Homo sapiens.

XX WO200009525-A2.

XX 24-FEB-2000.

XX 03-AUG-1999; 99WO-US17712.

XX 03-AUG-1998; 98US-0095212.

XX (UYEC-) UNIV EAST CAROLINA.

XX

PI Nyce JW;  
XX WPI; 2000-205971/18.  
XX New antisense oligonucleotides useful for treating e.g. pulmonary  
PT vasoconstriction, inflammation, allergies, asthma, hypertension,  
PT bronchitis, emphysema, respiratory distress syndrome, ischemia or  
PT cancers -  
XX Claim 18; Page 415; 1343pp; English.  
XX The present invention describes a new composition comprising an  
CC antisense oligonucleotide (ON) with low adenosine (up to 1%), which  
CC targets nucleic acids involved in bronchoconstriction, allergies, and/or  
CC inflammation. The ON can have antiinflammatory, antiallergic,  
CC antiasthmatic, cytostatic and analgesic activities. The compositions are  
CC useful for the treatment of diseases associated with inflammation,  
CC impaired airways, including lung disease and diseases whose secondary  
CC effects afflict the lungs of a subject. They can be used for treating  
CC e.g. ischaemic conditions, pulmonary vasoconstriction, allergies,  
CC asthma, impeded respiration, respiratory distress syndrome, pain, cystic  
CC fibrosis, pulmonary hypertension, emphysema, chronic obstructive  
CC pulmonary disease (COPD), and cancers such as leukaemias, lymphomas,  
CC carcinomas, and cancers which may metastasize to the lungs, including  
CC breast and prostate cancer. The reduction of the adenosine content of  
CC the ONs reduces side effects. The A-containing ONs break down with the  
CC release of deoxyadenosine which activates adenosine receptors causing  
CC bronchoconstriction and inflammation. AAA32313 to AAA3512 represent the  
CC nucleotide sequences given in the sequence listing from the present  
CC invention, which correspond to SEQ ID NO:1 to 2815, and then the last  
CC 185 sequences are also called SEQ ID NO:1 to 185, but the sequences  
CC differ from the previously named sequences. SEQ ID NO:11 to 1680  
CC (AAA32323 to AAA33992) are specifically claimed ONs from the present  
CC invention. N.B. Sequences given in the disclosure of the present  
CC invention do not match up with their corresponding SEQ ID NO: sequences  
XX given in the sequence listing.

XX Sequence 37 BP; 0 A; 11 C; 9 G; 17 T; 0 other;

Query Match 2.3%; Score 37; DB 21; Length 37;  
Best Local Similarity 100.0%; Pred. No. 1.4e+03;  
Matches 37; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 37 AGGACAAGAGCCAGGAGAAACACCGGAGGAGACCA 73  
Db 37 AGGACAAGAGCCAGGAGAAACACCGGAGGAGACCA 1

RESULT 5  
AAT76366/C

XX AAT76366 standard; DNA; 34 BP.

XX AAT76366;

XX 15-SEP-1997 (first entry)

XX Human interleukin 8 antisense oligonucleotide HUMTL8AAS3.

XX Asthma; airway epithelium; adenosine free; cystic fibrosis;  
KW chronic obstructive pulmonary disease; bronchitis; ss.

XX Synthetic.

XX WO9640162-A1.

XX 19-DEC-1996.

XX 06-JUN-1996; 96WO-US09306.

XX 07-JUN-1995; 95US-047497.

XX (UYEC-) UNIV EAST CAROLINA.

XX





PT Low adenosine (A) content antisense oligonucleotides which do not  
 PT trigger adenosine receptors during metabolism, useful e.g. for treating  
 PT cancers and respiratory obstructions -  
 XX  
 XX  
 XX  
 XX Claim 14; Page 236; 1592pp; English.

XX The present invention describes low adenosine (A) content antisense  
 CC oligonucleotides and compositions (I) comprising them. In the antisense  
 CC oligonucleotides the A is replaced by a 'Universal' or alternative base.  
 CC (I) can have respiratory, bronchodilator, antiinflammatory, analgesic,  
 CC immunosuppressive, antiasthmatic, hypotensive and cytostatic activities.  
 CC The antisense oligonucleotides and (I) can be used to down-regulate the  
 CC expression and/or activity of target polypeptides associated with  
 CC lung/respiratory disorders and malignancies, such as stimulating and  
 CC activating peptide factors and transmitters, transcription factors,  
 CC immunoglobulins and antibodies, antibody receptors, cytokines and  
 CC chemokines, endogenously produced specific and non-specific enzymes,  
 CC binding proteins, adhesion molecules and their receptors, cytokine and  
 CC chemokine receptors, adenosine receptors, bradykinin receptors, central  
 CC nervous system (CNS) and peripheral nervous and non-nervous system  
 CC receptors, CNS and peripheral nervous and non-nervous system peptide  
 CC transmitters, defensins, growth factors, vasoactive peptides and  
 CC receptors, binding proteins and malignancy associated proteins. The  
 CC antisense oligonucleotides may be used in this way to treat disorders  
 CC including respiratory obstruction (especially pulmonary obstruction  
 CC and/or bronchoconstriction) and/or lung inflammation, allergy(ies)  
 CC and/or surfactant hypoproduction which are associated with a disease or  
 CC condition selected from pulmonary vasoconstriction, inflammation,  
 CC allergies, asthma, impeded respiration, respiratory distress syndrome  
 CC (RDS), pain, cystic fibrosis (CF), allergic rhinitis (AR), pulmonary  
 CC hypertension, emphysema, chronic obstructive pulmonary disease (COPD),  
 CC pulmonary transplantation rejection, pulmonary infections, bronchitis,  
 CC and/or cancer. AAF18434 to AAF21543 represent human polynucleotide  
 CC fragments and antisense oligonucleotides used in the exemplification of  
 CC the present invention.

XX Sequence 34 BP; 0 A; 11 C; 7 G; 16 T; 0 other;

Query Match 2.1%; Score 34; DB 21; Length 34;  
 Best Local Similarity 100.0%; Pred. No. 4.5e+03;  
 Matches 34; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Cy 37 AGGACAAGAGCCAGGAAGAAACCCCGGAAGGAA 70  
 Db 34 AGGACAAGAGCCAGGAAGAAACCCCGGAAGGAA 1

RESULT 8  
 AAA33611/c  
 ID AAA33611 standard; DNA; 34 BP.

XX AC AAA33611;

XX 28-JUL-2000 (first entry)

DE Low adenosine antisense oligonucleotide SEQ ID NO:1300.

XX Human: adenosine receptor; low adenosine antisense oligonucleotide;  
 KW phosphorothioate; impaired respiration; inflammation; allergy;  
 KW allergic disease; bronchoconstriction; inhibitor; antiinflammatory;  
 KW antiasthmatic; cytostatic; analgesic; impaired airway;  
 KW lung disease; ischaemic condition; pulmonary vasoconstriction; asthma;  
 KW respiratory distress syndrome; pain; cystic fibrosis; emphysema;  
 KW pulmonary hypertension; chronic obstructive pulmonary disease; COPD;  
 KW cancer; leukaemia; lymphoma; carcinoma; metastasis; ss.

XX Homo sapiens.

XX WO200009525-A2.

XX 24-FEB-2000.

XX 03-AUG-1999; 99WO-US17712.

XX 03-AUG-1998; 98US-0095212.  
 XX (UYEC-) UNIV EAST CAROLINA.  
 PA Nyce JW;  
 PI WPI; 2000-205971/18.

XX New antisense oligonucleotides useful for treating e.g. pulmonary  
 CC vasoconstriction, inflammation, allergies, asthma, hypertension,  
 CC bronchitis, emphysema, respiratory distress syndrome, ischemia or  
 CC cancers -  
 XX Claim 18; Page 427; 1343pp; English.

XX The present invention describes a new composition comprising an  
 CC antisense oligonucleotide (ON) with low adenosine (up to 15%), which  
 CC targets nucleic acids involved in bronchoconstriction, allergies, and/or  
 CC inflammation. The ON can have antiinflammatory, antiasthmatic,  
 CC antiasthmatic, cytostatic and analgesic activities. The compositions are  
 CC useful for the treatment of diseases associated with inflammation,  
 CC impaired airways, including lung disease and diseases whose secondary  
 CC effects afflict the lungs of a subject. They can be used for treating  
 CC e.g. ischaemic conditions, pulmonary vasoconstriction, allergies,  
 CC asthma, impeded respiration, respiratory distress syndrome, pain, cystic  
 CC fibrosis, pulmonary hypertension, emphysema, chronic obstructive  
 CC pulmonary disease (COPD), and cancers such as leukaemias, lymphomas,  
 CC carcinomas, and cancers which may metastasize to the lungs, including  
 CC breast and prostate cancer. The reduction of the adenosine content of  
 CC the ONs reduces side effects. The A-containing ONs break down with the  
 CC release of deoxyadenosine which activates adenosine receptors causing the  
 CC bronchoconstriction and inflammation. AAA32313 to AAA35312 represent the  
 CC nucleotide sequences given in the sequence listing from the present  
 CC invention, which correspond to SEQ ID NO:1 to 2815, and then the last  
 CC 185 sequences are also called SEQ ID NO:1 to 185, but the sequences  
 CC differ from the previously named sequences. SEQ ID NO:11 to 1680  
 CC (AAA32323 to AAA33992) are specifically claimed ONs from the present  
 CC invention. N.B. Sequences given in the disclosure of the present  
 CC invention do not match up with their corresponding SEQ ID NO: sequences  
 CC given in the sequence listing.

XX Sequence 34 BP; 0 A; 11 C; 7 G; 16 T; 0 other;

Query Match 2.1%; Score 34; DB 21; Length 34;  
 Best Local Similarity 100.0%; Pred. No. 4.5e+03;  
 Matches 34; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Cy 37 AGGACAAGAGCCAGGAAGAAACCCCGGAAGGAA 70  
 Db 34 AGGACAAGAGCCAGGAAGAAACCCCGGAAGGAA 1

RESULT 9  
 AAA337164/c  
 ID AAA337164 standard; DNA; 33 BP.

XX AC AAA337164;

XX 23-JUN-1993 (first entry)

XX Anti-sense primer to amplify IL-8.

XX Interleukin-8; polymerase chain reaction; PCR;  
 KW cytokine synthesis inhibitor; inflammation;  
 KW monokine production; Southern analysis; ss.

XX Synthetic.

XX WO9302693-A.

XX 18-FEB-1993.

PF 06-AUG-1992; 92WO-US06378.  
 XX  
 PR 06-AUG-1991; 91US-0742129.  
 XX  
 PA (SCHE ) SCHERING CORP.  
 XX  
 PI De Waal Malefyt R, Howard M, Hsu DH, Ishida H, OGarra A;  
 PI Spits H, Zlotnik A;  
 XX  
 DR WPI: 1993-076172/09.  
 XX  
 XX Use of interleukin-10 to modulate inflammation or T-cell mediated  
 PT immune function - for treating septic and toxic shock,  
 PT auto-immune diseases, tumours and infectious diseases  
 XX  
 PS Example B6; Page 87; 208pp; English.  
 XX  
 CC To determine at which level IL-10 inhibited the production of  
 CC cytokines by monocytes, comparative PCR analyses were performed on  
 CC RNA isolated from monocytes, activated by LPS in the presence or  
 CC absence of IL-10, IL-4 or the neutralising anti-IL-10 MAb for 24  
 CC hours. mRNA isolated from the samples was reverse transcribed into  
 CC cDNA and amplified by cytokine-specific primers. Primer AAQ37164 is  
 CC specific to IL-8 (see AAQ37163 for sense primer). It was  
 CC found that IL-1 alpha, IL-6, TNF alpha, GM-CSF and G-CSF expression  
 CC was strongly inhibited by IL-10 and IL-4 at the mRNA level. IL-1 beta  
 CC and IL-8 expression was only slightly affected by IL-10.  
 XX  
 SQ Sequence 33 BP; 9 A; 10 C; 2 G; 12 T; 0 other;

Query Match 2.0%; Score 33; DB 14; Length 33;  
 - Best Local Similarity 100.0%; Pred. No. 6.5e+03;  
 Matches 33; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 358 GAGAGAGTTTTCAGAGGGCTGAGATTTCATAA 390  
 DB 33 GAGAGAGTTTTCAGAGGGCTGAGATTTCATAA 1

RESULT 10  
 AAT38979/c  
 ID AAT38979 standard; DNA; 33 BP.  
 XX  
 AC AAT38979;  
 XX  
 DT 29-MAY-1997 (first entry)  
 XX  
 DE Interleukin IL-8 3' PCR primer.  
 XX  
 KW Cytokine; expression profile; genital wart; interleukin 12; IL-12;  
 KW tumour regression; adjuvant; polymerase chain reaction; PCR;  
 KW condyloma acuminata; human papilloma virus; HPV-6; HPV-11; HPV16;  
 KW HPV18; anogenital; cutaneous; laryngeal; oesophageal; cancer; ss.  
 XX  
 OS Synthetic.  
 XX  
 PN WO9629091-Al.  
 XX  
 PD 26-SEP-1996.  
 XX  
 PF 22-MAR-1996; 96WO-GB00686.  
 XX  
 PR 22-MAR-1995; 95GB-0005784.  
 XX  
 PA (UYCA-) UNIV CAMBRIDGE TECH SERVICES LTD.  
 XX  
 PI Scarpini CG, Stanley MA;  
 XX  
 DR WPI: 1996-442947/44.  
 XX  
 PT Use of interleukin-12 to treat papilloma virus-associated lesions -  
 PT esp. as a vaccine adjuvant with papilloma virus antigen for  
 PT immuno-therapy of warts or tumours

XX  
 PS Disclosure; Page 14; 32pp; English.  
 XX  
 CC RNA was extracted from genital lesions, reverse transcribed to  
 CC produce cDNA and then the cDNA was used as the template for PCR  
 CC amplification of various cytokines using the primers in AAT38964-  
 CC AAT39005. To confirm the identity of amplified cDNA, digoxigenin-  
 CC labelled probes specific for each cytokine (see AAT39006-T39021)  
 CC were hybridised with Southern blots of amplified sequences. The  
 CC expression profile for regressing and non-regressing warts was  
 CC established and compared to cytokine expression patterns in normal  
 CC cervical tissue. Results showed that interleukin 12 is barely  
 CC expressed (if at all) in non-regressing warts, but is expressed in  
 CC regressing warts. This suggests a central role for IL-12 in wart  
 CC regression. It has been found that IL-12 can be used (especially  
 CC as a vaccine adjuvant) for treating papilloma virus-associated  
 CC lesions such as condyloma acuminata (anogenital warts) caused by  
 CC human papilloma virus type 6 (HPV-6) and/or HPV-11 and more  
 CC generally for treatment of tumours associated with HPV16 and HPV18  
 CC infection e.g. anogenital, cutaneous, laryngeal and oesophageal  
 CC cancers.  
 XX  
 SQ Sequence 33 BP; 9 A; 10 C; 2 G; 12 T; 0 other;

Query Match 2.0%; Score 33; DB 17; Length 33;  
 Best Local Similarity 100.0%; Pred. No. 6.5e+03;  
 Matches 33; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 358 GAGAGAGTTTTCAGAGGGCTGAGATTTCATAA 390  
 DB 33 GAGAGAGTTTTCAGAGGGCTGAGATTTCATAA 1  
 RESULT 11  
 AAV65360/c  
 ID AAV65360 standard; DNA; 33 BP.  
 XX  
 AC AAV65360;  
 XX  
 DT 22-JAN-1999 (first entry)  
 XX  
 DE Interleukin-8 antisense primer.  
 XX  
 KW Interleukin; IL-10; IL-6; IL-8; inflammatory response; endotoxin;  
 KW tumour necrosis factor alpha; TNF; IL-1 alpha; IL-1 beta; bacteria;  
 KW septic; toxic shock; superantigen; T-cell dependent response; human;  
 KW viral; PCR primer; ss.  
 XX  
 OS Synthetic.  
 OS Homo sapiens.  
 XX  
 PN US5837293-A.  
 XX  
 PD 17-NOV-1998.  
 XX  
 PF 07-JUN-1995; 95US-0481560.  
 XX  
 PR 24-MAR-1995; 95US-0410654.  
 PR 06-AUG-1991; 91US-0742129.  
 PR 06-AUG-1992; 92US-0926853.  
 PR 19-APR-1994; 94US-0229854.  
 PR 07-JUN-1995; 95US-0481560.  
 XX  
 PA (SCHE ) SCHERING CORP.  
 XX  
 PI De Waal Malefyt R, Howard M, Hsu D, Ishida H, O'Garra A;  
 PI Spits H, Zlotnik A;  
 XX  
 DR WPI: 1999-023391/02.  
 XX  
 PT Reducing inflammatory response by treatment with interleukin-10 -  
 PT particularly for treating bacterial infections, e.g. septic or toxic  
 PT shock



CC an agent coated on a microtitre plate is added to the PCR product. A  
 CC mixture of hybridised probe and PCR product, unhybridised probe and  
 CC PCR products and excess primers are transferred to a microtitre plate  
 CC coated with an agent capable of binding to a molecule on the PCR  
 CC product, or the molecule on the probe. The detectable marker is  
 CC detected on a bound hybridised probe and PCR product, preferably  
 CC via the detection of an antibody that detects the labelled primer  
 CC or probe, so that mRNA levels can be determined. primer and probe  
 CC sequences which can be used for the detection of interleukin-8  
 CC include the present sequence.

XX  
 SQ Sequence 31 BP; 11 A; 8 C; 9 G; 3 T; 0 other;  
 Query Match 1.9%; Score 31; DB 18; Length 31;  
 Best Local Similarity 100.0%; Pred. No. 1.4e+04;  
 Matches 31; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 119 TGGCAGCCTTCTGATTTCTGCAGCTCTGTG 149  
 |||||  
 Db 31 TGGCAGCCTTCTGATTTCTGCAGCTCTGTG 1

RESULT 14  
 AAI30870  
 ID AAI30870 standard; DNA; 31 BP.  
 AC AAI30870;  
 XX  
 XX 18-OCT-2001 (first entry)  
 DT  
 DE Human single nucleotide polymorphism (SNP) IL8.  
 XX-  
 KW Human; resequence; genotype; disease; forensic; paternity testing;  
 KW single nucleotide polymorphism; SNP; ss.  
 XX  
 OS Homo sapiens.  
 XX  
 FH Key Location/Qualifiers  
 FT Variation replace(16,C)  
 FT /\*tag= a  
 FT /standard\_name= "single nucleotide polymorphism"  
 XX  
 XX WO200166800-A2.  
 PN  
 XX 13-SEP-2001.  
 PD  
 XX 07-MAR-2001; 2001WO-US07268.  
 PF  
 XX 07-MAR-2000; 2000US-0187510.  
 PR 22-MAY-2000; 2000US-0206129.  
 XX  
 XX (WHED ) WHITEHEAD INST BIOMEDICAL RES.  
 PA  
 XX Cargill M, Ireland JS, Lander ES;  
 PI  
 XX WPI; 2001-522952/57.  
 DR  
 XX  
 XX Nucleic acid molecules from the human genome which include polymorphic  
 PT sites, useful in methods for predicting the presence, absence or  
 PT severity of a particular phenotype or disorder (e.g. diabetes)  
 PT associated with a particular genotype -  
 XX  
 XX Claim 1; Page 113; 145pp; English.  
 PS  
 XX The invention relates to the identification of nucleic acid molecules  
 CC (AAI29313-AAI31314) from the human genome which include polymorphic sites  
 CC which can predispose individuals to disease. Various genes from a number  
 CC of individuals were resequenced and single nucleotide polymorphisms  
 CC (SNPs) in these genes discovered. The method is useful for predicting the  
 CC presence, absence or severity of a particular phenotype or disorder (e.g.  
 CC diabetes) associated with a particular genotype. The nucleic acids  
 CC containing the polymorphic sites may be useful in forensics and paternity  
 CC testing.

XX  
 SQ Sequence 31 BP; 12 A; 8 C; 4 G; 7 T; 0 other;  
 Query Match 1.9%; Score 31; DB 22; Length 31;  
 Best Local Similarity 100.0%; Pred. No. 1.4e+04;  
 Matches 31; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 220 TTCCACCCCAATTTATCAAGAAGTGTGAG 250  
 |||||  
 Db 1 TTCCACCCCAATTTATCAAGAAGTGTGAG 31

RESULT 15  
 AAT45757  
 ID AAT45757 standard; DNA; 30 BP.  
 XX  
 XX AAT45757;  
 AC  
 XX 17-FEB-1997 (first entry)  
 DT  
 XX Human interleukin-8 gene probe.  
 DE  
 XX Polymerase chain reaction; PCR; interleukin; IL; cytokine;  
 KW growth factor; animal model; stem cell; haematopoiesis; CD34;  
 KW infection; HIV; human immunodeficiency virus; immunomodulator;  
 KW immortalise; bone marrow; stromal cell; engraftment; determination;  
 KW study; research; ss.  
 XX  
 OS Homo sapiens.  
 XX  
 XX WO9617627-A2.  
 PN  
 XX 13-JUN-1996.  
 PD  
 XX 08-DEC-1995; 95WO-US15986.  
 PF  
 XX 09-DEC-1994; 94US-0352957.  
 PR  
 XX (GART/) GARTNER S.  
 PA (HALL/) HALL E.  
 PA (KAUS/) KAUSHAL S.  
 PA (KESS/) KESSLER S.  
 PA (LRUS/) LA RUSSA V.  
 PA (MOSC/) MOSCA J D.  
 XX  
 XX Gartner S, Hall E, Kaushal S, Kessler S, La Russa V;  
 PI Mosca JD;  
 PI  
 XX WPI; 1996-286928/29.  
 DR  
 XX Animal models for human haematopoiesis - have en-grafted human or  
 PT primate stem cells in the presence of immortalised bone marrow  
 PT stromal cells  
 PT  
 XX Example 1; Page 15; 43pp; English.  
 PS  
 XX AAT45754-T45762 are probes used to determine whether or not the  
 CC cytokines IL-1, IL-6, IL-8, GM-CSF, G-CSF, M-CSF, TGF-alpha and  
 CC stem cell factor are expressed by a human bone marrow stromal  
 CC cell line, Lof(11-10). The cells were found to produce the cytokines  
 CC which support the growth of CD34+ stem cells. The Lof(11-10) cells  
 CC were injected into SCID mice (previously irradiated to provide an  
 CC internal space for CD34+ cells to populate). Five to seven days  
 CC after injection the mice were injected with 3 to 5 human CD34+ cells.  
 CC After 3 weeks, human CD34+ cells were found in the bone marrow of  
 CC the mice. The immortalised bone marrow stem cells create a human  
 CC microenvironment supplying human cytokines in the animals to provide  
 CC for the engraftment, maintenance and differentiation of CD34+ stem  
 CC cells. Animal models created by administering Lof(11-10) cells are  
 CC used to study and determine the effectiveness of therapies against  
 CC disease such as HIV infection. They can also be used to assay for  
 CC haematopoietic growth factors, immunomodulators and/or immune  
 CC toxins.

XX SQ Sequence 30 BP; 10 A; 5 C; 10 G; 5 T; 0 other;

Query Match 1.8%; Score 30; DB 17; Length 30;  
Best Local Similarity 100.0%; Pred. No. 2.1e+04;  
Matches 30; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 241 GAACGAGAGTGATGAGAGTGAGGACAC 270  
|||||  
Db 1 GAACGAGAGTGATGAGAGTGAGGACAC 30

RESULT 16  
AAV61828/c  
ID AAV61828 standard; DNA; 30 BP.

XX AC AAV61828;  
XX DT 15-JAN-1999 (first entry)  
XX PCR primer for human GISP coding sequence.  
XX Glucocorticoid inducible suppressor protein; interleukin-8; rheumatism;  
XX intracellular signal transmission inhibitor; IL-8 promoter; allergy;  
XX inflammatory disease; bronchial asthma; therapy; PCR primer; ss.  
XX Synthetic.  
XX Homo sapiens.  
XX WO9838213-Al.  
XX 03-SEP-1998.  
XX 27-FEB-1998; 98WO-JP00836.  
XX 28-FEB-1997; 97JP-0062008.  
XX (CYTO-) INST CYTOSIGNAL RES INC.  
XX Ohtsuka T, Yoshida H;  
XX WPI; 1998-481142/41.  
XX Protein inhibiting activation of interleukin-8 promoter in response  
XX to extracellular stimulus - for use in diseases involving  
XX interleukin-8 expression, such as inflammatory disorders and asthma  
XX Example 5; Page 56; 72pp; Japanese.  
XX This sequence represents a PCR primer for DNA encoding the human  
XX glucocorticoid inducible suppressor protein (GISP) of the invention. The  
XX GISP is an intracellular signal transmission inhibitor, and inhibits the  
XX activation of interleukin-8 (IL-8) promoter activity in response to a  
XX specific extracellular stimulus (especially by interleukin-1 beta). The  
XX invention can be used in the treatment and prevention of disorders in  
XX which IL-8 expression is involved, such as inflammatory diseases,  
XX bronchial asthma, allergy and rheumatism.

XX SQ Sequence 30 BP; 3 A; 7 C; 8 G; 12 T; 0 other;

Query Match 1.8%; Score 30; DB 19; Length 30;  
Best Local Similarity 100.0%; Pred. No. 2.1e+04;  
Matches 30; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 14 GACAGCAGACACACAGCTTCTAGGACAA 43  
|||||  
Db 30 GACAGCAGACACACAGCTTCTAGGACAA 1

RESULT 17  
AAV35562/c  
ID AAV35562 standard; DNA; 30 BP.

AC AAV35562;  
XX 24-SEP-1998 (first entry)  
XX Human interleukin 18 promoter PCR primer #2 from WO9822581 Example 1.  
XX Transcription repressor protein; inflammatory disease; regulator;  
XX interleukin 8; IL-8; rheumatoid arthritis; asthma; ringworm;  
XX skin inflammation; septic shock; gene therapy; MRC-5 SV1 TGI cell;  
XX PCR primer; ss.  
XX Synthetic.  
XX Homo sapiens.  
XX WO9822581-Al.  
XX 28-MAY-1998.  
XX 12-NOV-1997; 97WO-JP04127.  
XX 15-NOV-1996; 96JP-0305043.  
XX (CYTO-) INST CYTOSIGNAL RES INC.  
XX Nagasawa Y, Yoshida H;  
XX WPI; 1998-312467/27.  
XX Transcription repressor protein for treatment of inflammatory  
XX diseases - consists of transcription regulating protein having part  
XX deleted which is not involved in DNA binding  
XX Example 1; Page 12; 62pp; Japanese.  
XX The present sequence represents a PCR primer used in an example from the  
XX present invention. The present invention describes a protein which has  
XX transcription repressor activity and consists of a transcription  
XX regulator protein from which a part (not involved in DNA binding) has  
XX been deleted, or derived from such a protein by addition, deletion or  
XX substitution of one or more amino acid residues. In particular, the  
XX protein is one which inhibits the transcription of the interleukin-8  
XX (IL-8) gene. The protein is used for the treatment and prevention of  
XX disorders such as inflammatory diseases mediated by interleukin-8, e.g.  
XX rheumatoid arthritis, asthma, ringworm, skin inflammation, septic shock.  
XX The DNA encoding the protein may be used in gene therapy.

XX SQ Sequence 30 BP; 3 A; 7 C; 8 G; 12 T; 0 other;

Query Match 1.8%; Score 30; DB 19; Length 30;  
Best Local Similarity 100.0%; Pred. No. 2.1e+04;  
Matches 30; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 14 GACAGCAGACACACAGCTTCTAGGACAA 43  
|||||  
Db 30 GACAGCAGACACACAGCTTCTAGGACAA 1

RESULT 18  
AAV35781/c  
ID AAV35781 standard; DNA; 30 BP.

XX AC AAV35781;  
XX 22-SEP-1998 (first entry)  
XX PCR primer IL8P2 of the invention.  
XX Induce; apoptosis; guanine phosphoribosyltransferase; gpt; luciferase;  
XX cytokine; tumour necrosis factor; interleukin-1; inhibitory effect;  
XX intracellular signal transmission; PCR primer; ss.  
XX Synthetic.

PN W05822578-A1.  
 XX  
 PD 28-MAY-1998.  
 XX  
 XX 12-NOV-1997; 97WO-JP04126.  
 PF  
 XX 15-NOV-1996; 96JP-0305163.  
 PR  
 XX (CYTO-) INST CYTOSIGNAL RES INC.  
 PA  
 XX Nagasawa Y, Yoshida H;  
 PI  
 XX WPI; 1998-312464/27.  
 DR  
 XX  
 XX Test system for detecting intra-cellular signal transmission  
 PT inhibition - using vector containing apoptosis-inhibiting or  
 PT reporter gene and promoter sequence, used for, e.g. screening for  
 PT potential anti-inflammatory agents  
 PT  
 XX Example 1; Page 20; 62pp; Japanese.  
 PS  
 XX PCR primers AAV35780-81 are used in the course of the invention. The  
 CC specification describes plasmid vectors which comprise a gene which  
 CC can induce apoptosis under specific conditions, e.g. guanine  
 CC phosphoribosyltransferase (gpt), or a reporter gene, e.g. luciferase,  
 CC where the gene is situated downstream of a promoter which responds to  
 CC specific extracellular stimulation such as the presence of a cytokine,  
 CC e.g. tumour necrosis factor (TNF) or interleukin-1. The vector may be  
 CC used to transform a suitable cell line, such as a cell line which does  
 CC not produce hypoxanthine-guanine phosphoribosyl transferase (HGPRT). The  
 CC transformed cells are used to test the inhibitory effect of a gene or  
 CC substance on intracellular signal transmission.  
 XX  
 XX Sequence 30 BP; 3 A; 7 C; 8 G; 12 T; 0 other;  
 SQ  
 Query Match 1.8%; Score 30; DB 19; Length 30;  
 Best Local Similarity 100.0%; Pred. No. 2.1e+04;  
 Matches 30; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 14 GACAGCAGAGCAGACAAAGCTTCTAGGACAA 43  
 DB 30 GACAGCAGAGCAGACAAAGCTTCTAGGACAA 1  
 RESULT 19  
 AAC88174/c  
 ID AAC88174 standard; DNA; 30 BP.  
 XX  
 AC AAC88174;  
 XX  
 DT 14-MAR-2001 (first entry)  
 XX  
 XX Human interleukin 8 5'-regulatory sequence PCR primer SEQ ID NO:3.  
 DE  
 XX Human; interleukin 8; IL-8; regulation; point mutation; NF-kappa B;  
 KW AP-1 binding site; interleukin 1 beta; tumour necrosis factor alpha;  
 KW promoter; antiinflammatory; antiarteriosclerotic; screening; inhibitor;  
 KW identification; inflammation; arteriosclerosis; PCR primer; ss.  
 XX  
 OS Homo sapiens.  
 XX  
 XX W0200071697-A1.  
 PN  
 XX 30-NOV-2000.  
 PD  
 XX 24-MAY-2000; 2000WO-JP03326.  
 PF  
 XX 24-MAY-1999; 99JP-0143032.  
 PR  
 XX (SANY ) SANKYO CO LTD.  
 PA  
 XX Koishi R, Yoshimura C, Serizawa N;  
 XX  
 XX WPI; 2002-049698/06.  
 PI  
 XX Lyamichiev V, Allawi H, Dong F, Neri BP, Vener IT;  
 XX  
 XX Identifying oligonucleotides hybridizing to nucleic acids containing  
 PT secondary structure, useful in clinical diagnosis, comprises  
 PT identifying primers that interact with the target to form an extension  
 PT product under amplification conditions -  
 XX  
 PS Claim 48; Fig 81A; 409pp; English.  
 XX  
 XX The present invention describes a method for identifying oligonucleotides

DR WPI; 2001-112080/12.  
 XX  
 XX Screening for potential inhibitors of the NF-kappa-B activation  
 PT pathway, useful for treatment of inflammation and arteriosclerosis -  
 PT  
 XX Example 1; Page 15; 49pp; Japanese.  
 PS  
 XX The present invention describes a method for screening for potential  
 CC specific inhibitors of pathways activated by interleukin (IL)-1 beta,  
 CC tumour necrosis factor (TNF) alpha or a leustrodaxin derivative, by  
 CC culturing a cell transformed by a marker gene controlled by a promoter  
 CC gene containing a NF-kappa-B binding site in the presence of  
 CC IL1beta/TNFalpha/leustrodaxin derivative and the inhibitor and observing  
 CC the degree of marker gene expression. Also described is a method for  
 CC activating a promoter gene containing a NF-kappa-B binding site and  
 CC inhibitors identified by the above method. The method can be used for  
 CC the identification of inhibitors of the NF-kappa-B activation pathway  
 CC which can be used for the treatment of inflammation and arteriosclerosis.  
 CC The present sequence represents a PCR primer for the 5'-regulatory  
 CC sequence of the human IL-8 gene, which is used in an example from the  
 CC present invention.  
 XX  
 XX Sequence 30 BP; 3 A; 7 C; 8 G; 12 T; 0 other;  
 SQ  
 Query Match 1.8%; Score 30; DB 22; Length 30;  
 Best Local Similarity 100.0%; Pred. No. 2.1e+04;  
 Matches 30; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 14 GACAGCAGAGCAGACAAAGCTTCTAGGACAA 43  
 DB 30 GACAGCAGAGCAGACAAAGCTTCTAGGACAA 1  
 RESULT 20  
 ABL46338/c  
 ID ABL46338 standard; DNA; 30 BP.  
 XX  
 AC ABL46338;  
 XX  
 DT 26-APR-2002 (first entry)  
 XX  
 XX Human interleukin-1 beta oligonucleotide SEQ ID NO:305.  
 DE  
 XX Nucleic acid accessible hybridisation site; detection; hybridisation;  
 KW characterisation; identification; nucleic acid structure; diagnosis;  
 KW PCR primer; probe; ss.  
 XX  
 OS Homo sapiens.  
 OS Synthetic.  
 XX  
 XX W0200198537-A2.  
 PN  
 XX 27-DEC-2001.  
 PD  
 XX 15-JUN-2001; 2001WO-US19401.  
 PF  
 XX 17-JUN-2000; 2000US-212308P.  
 PR  
 XX 15-JUN-2001; 2001US-0212308.  
 XX  
 XX (THIR-) THIRD WAVE TECHNOLOGIES INC.  
 PA  
 XX  
 XX Lyamichiev V, Allawi H, Dong F, Neri BP, Vener IT;  
 PI  
 XX WPI; 2002-049698/06.  
 DR  
 XX Identifying oligonucleotides hybridizing to nucleic acids containing  
 PT secondary structure, useful in clinical diagnosis, comprises  
 PT identifying primers that interact with the target to form an extension  
 PT product under amplification conditions -  
 XX  
 PS Claim 48; Fig 81A; 409pp; English.  
 XX  
 XX The present invention describes a method for identifying oligonucleotides

CC with desired hybridisation properties to nucleic acid targets containing  
CC secondary structure. The method comprises amplifying a target nucleic  
CC acid having at least one accessible and one inaccessible site. Primers  
CC that form an extension product are identified as the oligonucleotides  
CC which can interact with the folded target nucleic acid. Oligonucleotides  
CC from the present invention can be used in novel detection methods for  
CC clinical diagnostic purposes, including the detection and identification  
CC of pathogenic organisms (e.g. HIV). The method allows the ability to  
CC rapidly analyse nucleic acid structures. ABL46034 to ABL46367 represent  
CC sequences used in the exemplification of the present invention.

XX  
SQ Sequence 30 BP; 13 A; 4 C; 2 G; 11 T; 0 other;

Query Match 1.8%; Score 30; DB 24; Length 30;  
Best Local Similarity 100.0%; Pred. No. 2.1e+04;  
Matches 30; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1165 ATGATGTTTTATTAGATAAAATTTCAATCAG 1194

Db 30 ATGATGTTTTATTAGATAAAATTTCAATCAG 1

Search completed: February 9, 2003, 17:30:46  
Job time : 350 secs

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GenCore version 5.1.3  
Copyright (c) 1993 - 2003 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: February 9, 2003, 15:52:56 ; Search time 4173 Seconds  
(without alignments)  
11430.505 Million cell updates/sec

Title: US-09-960-143-3  
Perfect score: 1639  
Sequence: 1 acaacttcagagacagca.....atataattgtgtcacaagt 1639

Scoring table: IDENTITY\_NUC  
Gapop 10.0 , Gapext 1.0

Searched: 2054640 seqs, 14551402878 residues  
Total number of hits satisfying chosen parameters: 841850

Minimum DB seq length: 0  
Maximum DB seq length: 50

Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

Database : GenEmbl.\*

- 1: gb\_ba.\*
- 2: gb\_htg.\*
- 3: gb\_in.\*
- 4: gb\_om.\*
- 5: gb\_ov.\*
- 6: gb\_pat.\*
- 7: gb\_ph.\*
- 8: gb\_pl.\*
- 9: gb\_pr.\*
- 10: gb\_ro.\*
- 11: gb\_sts.\*
- 12: gb\_sy.\*
- 13: gb\_un.\*
- 14: gb\_vl.\*
- 15: em\_ba.\*
- 16: em\_fun.\*
- 17: em\_hum.\*
- 18: em\_in.\*
- 19: em\_mu.\*
- 20: em\_om.\*
- 21: em\_or.\*
- 22: em\_ov.\*
- 23: em\_pat.\*
- 24: em\_ph.\*
- 25: em\_pl.\*
- 26: em\_ro.\*
- 27: em\_sts.\*
- 28: em\_un.\*
- 29: em\_vl.\*
- 30: em\_htg\_hum.\*
- 31: em\_htg\_inv.\*
- 32: em\_htg\_other.\*
- 33: em\_htg\_mus.\*
- 34: em\_htg\_pln.\*
- 35: em\_htg\_rod.\*
- 36: em\_htg\_mam.\*
- 37: em\_htg\_vrt.\*
- 38: em\_sy.\*
- 39: em\_htgo\_hum.\*
- 40: em\_htgo\_mus.\*
- 41: em\_htgo\_other.\*

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,  
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match %	Length	DB	ID	Description
1	37.4	2.3	47	6	AR021184	AR021184 Sequence
2	37.4	2.3	47	6	I60565	I60565 Sequence 42
C 3	37	2.3	47	6	AR021182	AR021182 Sequence
C 4	37	2.3	37	6	I60563	I60563 Sequence 40
C 5	33	2.0	33	6	A56958	A56958 Sequence 16
C 6	33	2.0	33	6	AR052918	AR052918 Sequence
C 7	33	2.0	33	6	AR054281	AR054281 Sequence
C 8	33	2.0	33	6	AR054483	AR054483 Sequence
C 9	30.6	1.9	31	6	AX249279	AX249279 Sequence
C 10	30	1.8	30	6	AR200024	AR200024 Sequence
C 11	30	1.8	30	6	AX419968	AX419968 Sequence
C 12	30	1.8	30	6	BD004257	BD004257 Method fo
C 13	29	1.8	29	6	AX317616	AX317616 Sequence
C 14	28	1.7	28	6	AR090359	AR090359 Sequence
C 15	28	1.7	28	6	AR090360	AR090360 Sequence
C 16	28	1.7	28	6	AR197394	AR197394 Sequence
C 17	28	1.7	28	6	AR197395	AR197395 Sequence
C 18	26	1.6	26	6	AX280042	AX280042 Sequence
C 19	25	1.5	25	6	I22270	I22270 Sequence 8
C 20	25	1.5	26	6	AX280041	AX280041 Sequence
C 21	24	1.5	24	6	AR207732	AR207732 Sequence
C 22	23	1.4	23	6	I22269	I22269 Sequence 7
C 23	23	1.4	40	6	AR021188	AR021188 Sequence
C 24	22.8	1.4	31	6	AX203829	AX203829 Sequence
C 25	22.8	1.4	48	6	AX426642	AX426642 Sequence
C 26	22.6	1.4	48	6	AX426920	AX426920 Sequence
C 27	22.2	1.4	28	6	AX203828	AX203828 Sequence
C 28	22	1.3	22	6	AR130451	AR130451 Sequence
C 29	22	1.3	22	6	E09226	E09226 Primer for
C 30	22	1.3	22	6	E09227	E09227 Primer for
C 31	22	1.3	48	6	I33522	I33522 Sequence 1
C 32	22	1.3	50	8	CNS018YM	AL110965 Botrytis
C 33	21.8	1.3	47	6	AX114372	AX114372 Sequence
C 34	21.8	1.3	48	6	I33523	I33523 Sequence 2
C 35	21.6	1.3	41	6	AR164944	AR164944 Sequence
C 36	21.6	1.3	41	6	E22883	E22883 DNA sequence
C 37	21.6	1.3	50	6	AX160081	AX160081 Sequence
C 38	21.6	1.3	50	6	AX160578	AX160578 Sequence
C 39	21.4	1.3	43	6	AX484483	AX484483 Sequence
C 40	21.4	1.3	47	1	LACDRRH	M84772 Lactococcus
C 41	21.4	1.3	48	9	S64862S2	S64863 alpha 1-the
C 42	21.2	1.3	43	6	AX484516	AX484516 Sequence
C 43	21.2	1.3	45	8	AB008098	AB008098 Saccharom
C 44	21.2	1.3	50	6	AX157057	AX157057 Sequence
C 45	21	1.3	21	6	A56957	A56957 Sequence 15

ALIGNMENTS

RESULT 1  
AR021184  
LOCUS AR021184 47 bp DNA linear PAT 05-DEC-1998  
DEFINITION Sequence 42 from patent US 5789539.  
ACCESSION AR021184  
VERSION AR021184.1 GI:3975799  
KEYWORDS  
SOURCE Unknown.  
ORGANISM Unknown.  
REFERENCE Unclassified.  
1 (bases 1 to 47)  
AUTHORS Daly,T.J. and Larosa,G.J.  
TITLE Chemokine-like proteins and methods of use  
JOURNAL Patent: US 5789539-A 42 04-AUG-1998;  
FEATURES Location/Qualifiers

## RESULT 4

TITLE Use of interleukin-10 (IL-10) to treat endotoxin- or superantigen-induced toxicity

JOURNAL Patent: US 5837293-A 42 10-NOV-1998;

FEATURES Location/Qualifiers

Source

BASE COUNT 9 a 10 c 2 g 12 t

ORIGIN

Query Match

Best Local Similarity 2.0%; Score 33; DB 6; Length 33;

Matches 33; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 358 GAGAAGTTTTGAAGAGGGCTGAGAATTCATAA 390

Db 33 GAGAAGTTTTGAAGAGGGCTGAGAATTCATAA 1

RESULT 7

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

FEATURES

Source

BASE COUNT

ORIGIN

Query Match

Best Local Similarity

Matches

QY 358

Db 33

RESULT 8

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

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REFERENCE

AUTHORS

TITLE

JOURNAL

FEATURES

Source

BASE COUNT

ORIGIN

Query Match

Best Local Similarity

Matches

QY 14

Db 30

RESULT 11

LOCUS

DEFINITION

ACCESSION

Sequence 1358 from Patent WO0166800.

Version AX249279.1 GI:15863902

Keywords human.

Organism Homo sapiens

Reference 1 (bases 1 to 31)

Authors Cargill, M., Ireland, J. S. and Lander, E. S.

Title Human single nucleotide polymorphisms

Journal Patent: WO 0166800-A 1358 13-SEP-2001;

Features WHITEHEAD INSTITUTE FOR BIOMEDICAL RESEARCH (US)

Location/Qualifiers

Source 1.31

Base Count 11 a 8 c 4 g 7 t

Origin 1 others

Query Match

Best Local Similarity

Matches

QY 220

Db 1

RESULT 10

LOCUS

DEFINITION

ACCESSION

Version AR2000024.1 GI:20250098

Keywords Unknown.

Organism Unclassified.

Reference 1 (bases 1 to 30)

Authors Negasawa, Y. and Yoshida, H.

Title Transcriptional inhibitor

Journal Patent: US 6355775-A 6 12-MAR-2002;

Features Location/Qualifiers

Source 1.30

Base Count 3 a 7 c 8 g 12 t

Origin 12 t

Query Match

Best Local Similarity

Matches

QY 14

Db 30

RESULT 11

LOCUS

DEFINITION

ACCESSION

Sequence 305 from Patent WO0198537.

Version AX419968

Keywords human.

Organism Homo sapiens

Reference 1 (bases 1 to 30)

Authors Negasawa, Y. and Yoshida, H.

Title Transcriptional inhibitor

Journal Patent: US 6355775-A 6 12-MAR-2002;

Features Location/Qualifiers

Source 1.30

Base Count 3 a 7 c 8 g 12 t

Origin 12 t

Query Match

Best Local Similarity

Matches

QY 14

Db 30

RESULT 11

LOCUS

DEFINITION

ACCESSION

Sequence 305 from Patent WO0198537.

Version AX419968

Keywords human.

Organism Homo sapiens

Reference 1 (bases 1 to 30)

Authors Negasawa, Y. and Yoshida, H.

Title Transcriptional inhibitor

Journal Patent: US 6355775-A 6 12-MAR-2002;

Features Location/Qualifiers

Source 1.30

Base Count 3 a 7 c 8 g 12 t

Origin 12 t

—2—

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LOCUS       AX280042                26 bp      DNA      linear      PAT 02-NOV-2001
DEFINITION   Sequence 17 from Patent WO0177382.
ACCESSION    AX280042
VERSION      AX280042.1  GI:16607493
KEYWORDS     synthetic construct.
SOURCE       synthetic construct
ORGANISM     artificial sequences.
REFERENCE    1
AUTHORS      Hull, J. and Kwiatkowski, D.P.
TITLE        Genetic factor affecting cytokine expression
JOURNAL      Patent: WO 0177382-A 17 18-OCT-2001;
             ISIS INNOVATION LIMITED (GB)
FEATURES     Location/Qualifiers
             1..26
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Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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Db 1 CCCAGTTAAATTTTCATTTTCAGATAA 26
102/153
RESULT 19
LOCUS       I22270/c                25 bp      DNA      linear      PAT 07-OCT-1996
DEFINITION   Sequence 8 from patent US 5527678.
ACCESSION    I22270
VERSION      I22270.1  GI:1602624
KEYWORDS     Unknown.
SOURCE       Unknown.
ORGANISM     Unclassified.
REFERENCE    1 (bases 1 to 25)
AUTHORS      Blaser, M.J., Tummuru, M.K.R. and Sharma, S.A.
TITLE        CagB and CagC genes of helicobacter pylori and related compositions
JOURNAL      Patent: US 5527678-A 8 18-JUN-1996;
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Best Local Similarity 100.0%; Pred. No. 2.6e+06;
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QY 358 GAGAAGTTTTTGAAGAGGCGTGAGA 382
Db 25 GAGAAGTTTTTGAAGAGGCGTGAGA 1
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LOCUS       AX280041                26 bp      DNA      linear      PAT 02-NOV-2001
DEFINITION   Sequence 16 from Patent WO0177382.
ACCESSION    AX280041
VERSION      AX280041.1  GI:16607492
KEYWORDS     synthetic construct.
SOURCE       synthetic construct
ORGANISM     artificial sequences.
REFERENCE    1
AUTHORS      Hull, J. and Kwiatkowski, D.P.
TITLE        Genetic factor affecting cytokine expression
JOURNAL      Patent: WO 0177382-A 16 18-OCT-2001;

LOCUS       AX280042                7 t
DEFINITION   Sequence 17 from Patent WO0177382.
ACCESSION    AX280042
VERSION      AX280042.1  GI:16607493
KEYWORDS     synthetic construct.
SOURCE       synthetic construct
ORGANISM     artificial sequences.
REFERENCE    1
AUTHORS      Hull, J. and Kwiatkowski, D.P.
TITLE        Genetic factor affecting cytokine expression
JOURNAL      Patent: WO 0177382-A 17 18-OCT-2001;
             ISIS INNOVATION LIMITED (GB)
FEATURES     Location/Qualifiers
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             /organism="synthetic construct"
             /db_xref="taxon:32630"
             /note="Oligonucleotide"
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Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 248 GAGTGATTGAGAGTGACACACTGGCG 275
Db 28 GAGTGATTGAGAGTGACACACTGGCG 1
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LOCUS       AR197394                28 bp      DNA      linear      PAT 20-APR-2002
DEFINITION   Sequence 479 from patent US 6352829.
ACCESSION    AR197394
VERSION      AR197394.1  GI:20247243
KEYWORDS     Unknown.
SOURCE       Unknown.
ORGANISM     Unclassified.
REFERENCE    1 (bases 1 to 28)
AUTHORS      Chenchik, A., Jokhadze, G. and Bibilashvili, R.
TITLE        Methods of assaying differential expression
JOURNAL      Patent: US 6352829-A 479 05-MAR-2002;
FEATURES     Location/Qualifiers
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             /organism="unknown"
BASE COUNT     6 a 9 c 7 g 6 t
ORIGIN
Query Match      1.7%; Score 28; DB 6; Length 28;
Best Local Similarity 100.0%; Pred. No. 8.8e+05;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 88 AACATGACTTCCAAGTGCGCGTGCTC 115
Db 1 AACATGACTTCCAAGTGCGCGTGCTC 28
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LOCUS       AR197395/c                28 bp      DNA      linear      PAT 20-APR-2002
DEFINITION   Sequence 480 from patent US 6352829.
ACCESSION    AR197395
VERSION      AR197395.1  GI:20247244
KEYWORDS     Unknown.
SOURCE       Unknown.
ORGANISM     Unclassified.
REFERENCE    1 (bases 1 to 28)
AUTHORS      Chenchik, A., Jokhadze, G. and Bibilashvili, R.
TITLE        Methods of assaying differential expression
JOURNAL      Patent: US 6352829-A 480 05-MAR-2002;
FEATURES     Location/Qualifiers
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             /organism="unknown"
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Best Local Similarity 100.0%; Pred. No. 8.8e+05;
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QY 248 GAGTGATTGAGAGTGACACACTGGCG 275
Db 28 GAGTGATTGAGAGTGACACACTGGCG 1
RESULT 18
LOCUS       AX280042                26 bp      DNA      linear      PAT 02-NOV-2001
DEFINITION   Sequence 16 from Patent WO0177382.
ACCESSION    AX280042
VERSION      AX280042.1  GI:16607492
KEYWORDS     synthetic construct.
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AUTHORS      Hull, J. and Kwiatkowski, D.P.
TITLE        Genetic factor affecting cytokine expression
JOURNAL      Patent: WO 0177382-A 16 18-OCT-2001;

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